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Number 1 of 1

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U.S. PROVISIONAL PATENT APPLICATION

ENTITLED:

2,4,5-Trisubstituted Imidazoles and Their Use
as Anti-Cancer Agents

Inventor(s):

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CERTIFICATE OF EXPRESS MAILING

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2,4,5-TRISUBSTITUTED IMIDAZOLES AND THEIR USE AS ANTI-CANCER AGENTS

FIELD OF INVENTION

5 This invention pertains to the field of anti-cancer compounds and, in particular, to the use of therapeutically active 2,4,5-trisubstituted imidazole compounds in the treatment of cancer.

BACKGROUND OF THE INVENTION

10 A cancer is a malignant tumour of potentially unlimited growth. It is primarily the pathogenic replication (a loss of normal regulatory control) of various given types of cells found in the human body. By select mutation resulting from a primary lesion, the DNA of a cancer cell evolves and converts the cell into an autonomous system. Conventional cancer treatments have focused mainly on killing cancerous cells. Chemotherapeutic agents currently used for anti-cancer/anti-tumour therapy are selected for their toxicity towards rapidly proliferating cells. Most of them cause undesirable systemic effects such as cardiac or renal toxicity, marrow aplasia, alopecia, nausea and vomiting. During the last few years, many researchers have tried to eliminate these side effects by developing drugs having suitable physico-chemical properties allowing an increase of the availability of the drug to the tumour site. New molecules extracted from natural sources, synthetically or semi-synthetically produced, enzymes, radioisotopes, DNA toxins, various macromolecules, and antibodies against fibrin or against tumour-specific surface antigens are bound to drugs in an attempt to increase selectivity of the chemotherapeutic agents.

20 The effectiveness of most anticancer agents is greatly reduced because of their high toxicity and the nature of the illness. It is believed that the problem of high toxicity of the anticancer agents can be circumvented by chemical modifications of those structures in such a way that they act more specifically on tumour cells without increasing systemic toxicity. The research in this field is therefore mainly directed to the synthesis of anticancer agents which would possess high antineoplastic activity, low systemic toxicity and low mutagenicity on normal cells.

25

Heterocyclic compounds, especially heterocyclic azole derivatives, have been shown to have a wide spectrum of biological activities. One class of compounds with interesting biological activities is the imidazoles (derivatives containing a five-membered heterocyclic azole). A variety of biological activities have been reported for imidazole derivatives with different substitution patterns (Lee *et al. Nature* 1994 327:739-745; Abdel-Meguid *et al. Biochemistry*, 1994, 33:11671; Heerding *et al. Bioorg. Med. Chem. Lett.* 2001, 11:2061-2065; Bu *et al. Tetrahedron Lett.* 1996, 37:7331-7334; Lewis JR. *Nat. Prod. Rep.* 1999, 16:389-418; Lewis JR. *Nat. Prod. Rep.* 1998, 15:417-437 and 371-395).

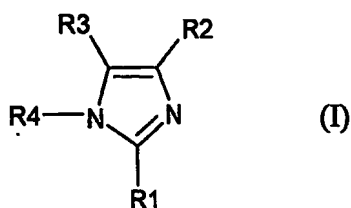
Biological activities have also been reported for aryl-imidazole derivatives, for example, these compounds can act as modulators of multi-drug resistance in cancer cells (Zhang *et al. Bioorg. Med. Chem. Lett.* 2000, 10:2603-2605), inhibitors of p38 MAP kinase (Adams *et al. Bioorg. Med. Chem. Lett.* 2001, 11:867-2870, McLay *et al. Bioorg. Med. Chem.* 2001, 9:537-554) and of cytokines (U.S. Patent Nos. 5,656,644; 5,686,455; 5,916,891; 5,945,418; and 6,268,370), and inhibitors of bacterial growth (Antolini *et al. Bioorg. Med. Chem. Lett.* 1999, 9:1023-1028).

A few reports have indicated that triaryl-imidazole compounds can act as inhibitors of p38 MAP kinase (for example, see LoGrasso *et al. Biochemistry.* 1997, 36:10422-10427) and as modulators of multi-drug resistance in cancer cells (Sarshar *et al. Bioorg. Med. Chem. Lett.* 2000, 10:2599-2601), however, the majority of the literature indicates that these compounds have found use mainly as colour producing reagents (U.S. Patent Nos. 4,089,747; 5,024,935; 5,047,318; 5,496,702; 5,514,550; and 5,693,589) and as photopolymerization initiators (U.S. Patent Nos. 6,117,609 and 6,060,216), generally in dimeric form.

This background information is provided for the purpose of making known information believed by the applicant to be of possible relevance to the present invention. No admission is necessarily intended, nor should be construed, that any of the preceding information constitutes prior art against the present invention.

SUMMARY OF THE INVENTION

It is an object of the present invention to provide a class of compounds which are 2,4,5-trisubstituted imidazole derivatives that have anti-cancer activity. In accordance with an aspect of the present invention there is provided a use of a compound having structural formula (I), or a salt thereof, as an anti-cancer agent:



wherein:

R1 is aryl, substituted aryl, heterocycle, substituted heterocycle, heteroaryl, or substituted heteroaryl;

R2 and R3 are independently aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, or substituted heteroaryl or R2 and R3 when taken together along with the carbon atoms they are attached to, form aryl or substituted aryl, heterocycle, substituted heterocycle, heteroaryl, or substituted heteroaryl and

R4 is hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano.

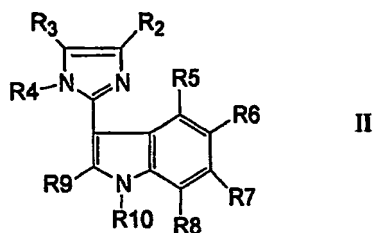
In accordance with another aspect of the present invention, there is provided a use of a compound having structural formula (I), or a salt thereof, in the preparation of an anti-cancer composition.

In accordance with another aspect of the present invention, there is provided a method of inhibiting the growth and/or proliferation of neoplastic cells comprising contacting said cells with an effective amount of a compound having general formula (I), or a salt thereof.

In accordance with another aspect of the present invention there is provided a method of inhibiting neoplastic cell growth in a mammal comprising administering to said mammal a therapeutically effective amount of a compound having general formula (I), or a salt thereof.

5 In accordance with another aspect of the present invention, there is provided an anti-cancer composition comprising an effective amount of a compound having structural formula (I), or a salt thereof, and a carrier, diluent or excipient.

In accordance with another aspect of the present invention, there is provided a compound having the structural formula:



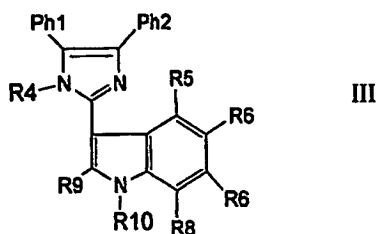
10 or a salt thereof, wherein:

R2 and R3 are independently aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, or substituted heteroaryl or R2 and R3 when taken together along with the carbon atoms they are attached form aryl or substituted aryl, heterocycle, heteroaryl, substituted heterocycle, or substituted heteroaryl;

15 R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or
20 cyano;

R10 is H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, acyl, -CH₂-aryl, -CH₂-heteroaryl.

In accordance with another aspect of the present invention, there is provided a compound having the structural formula:

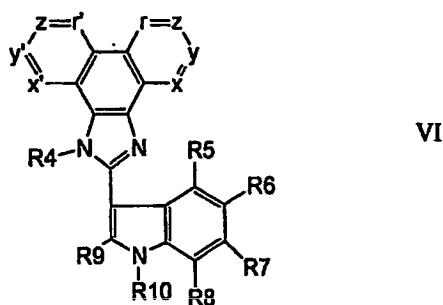


or a salt thereof, wherein:

Ph1 and Ph2 are independently selected from phenyl and substituted phenyl;

- 5 R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or
- 10 cyano;
- R10 is H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, acyl, -CH₂-aryl, -CH₂-heteroaryl.

- In accordance with another aspect of the present invention, there is provided a compound having
- 15 the structural formula:



or a salt thereof, wherein:

- R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino,
- 20

amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano;

R10 is H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, acyl, -CH₂-aryl, -CH₂-heteroaryl;

x is CR11 or N;

y is CR12 or N;

z is CR13 or N;

r is CR14 or N;

10 x' is CR15 or N;

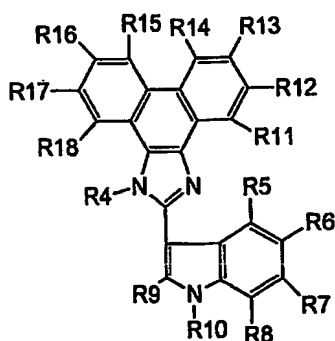
y' is CR16 or N;

z' is CR17 or N;

x' is CR18 or N;

15 R11, R12, R13, R14, R15, R16, R17 and R18 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, alkenyl, alkenyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano.

20 In accordance with another aspect of the present invention, there is provided a compound having the structural formula:



VII

or a salt thereof, wherein:

R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl,

substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano;

- 5 R10 is H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, acyl, -CH₂-aryl, -CH₂-heteroaryl;
R11, R12, R13, R14, R15, R16, R17 and R18 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, alkenyl, alkenyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl,
10 heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano.

In accordance with another aspect of the present invention, there is provided a method of inhibiting the growth and/or proliferation of neoplastic cells comprising contacting said cells with an effective amount of a compound selected from the compounds of general formula (II),
15 (III), (IV), (V), (VI) and (VII) or a salt thereof.

In accordance with another aspect of the present invention there is provided a method of inhibiting neoplastic cell growth in a mammal comprising administering to said mammal a therapeutically effective amount of a compound selected from the compounds of general formula (II), (III), (IV), (V), (VI) and (VII) or a salt thereof.

20

BRIEF DESCRIPTION OF THE FIGURES

Figure 1 depicts the effect of compounds of Formula I on growth of human adenocarcinoma HT29 cells in CD-1 nude mice.

Figure 2 depicts the effects of compounds of Formula I on the average weight of tumours in CD-1 nude mice (average weight per group of mice).

- 25 **Figure 3** depicts the effects of compounds of Formula I on the weight of tumours in CD-1 nude mice (individual tumour weights).

DETAILED DESCRIPTION OF THE INVENTION

The present invention provides a class of 2,4,5-trisubstituted imidazole compounds and for their use as anti-cancer agents. The present invention further provides for methods of inhibiting neoplastic cell growth and/or proliferation in an animal by administering to the animal an effective amount of a compound of Formula I, either alone or in combination with one or more standard chemotherapeutics.

Definitions

Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention pertains.

The terms are defined as follows:

The term "halogen" refers to fluorine, bromine, chlorine, and iodine atoms.

The term "hydroxyl" refers to the group -OH.

The term "thiol" or "mercapto" refers to the group -SH, and -S(O)₀₋₂.

The term "lower alkyl" refers to a straight chain or branched, or cyclic, alkyl group of one to ten carbon atoms. This term is further exemplified by such groups as methyl, ethyl, *n*-propyl, *i*-propyl, *n*-butyl, *t*-butyl, *i*-butyl (or 2-methylpropyl), cyclopropylmethyl, *i*-amyl, *n*-amyl, hexyl and the like.

The term "substituted lower alkyl" refers to lower alkyl as just described including one or more groups such as hydroxyl, thiol, alkylthiol, halogen, alkoxy, amino, amido, carboxyl, cycloalkyl, substituted cycloalkyl, heterocycle, cycloheteroalkyl, substituted cycloheteroalkyl, acyl, carboxyl, aryl, substituted aryl, aryloxy, hetaryl, substituted hetaryl, aralkyl, heteroaralkyl, alkyl alkenyl, alkyl alkynyl, alkyl cycloalkyl, alkyl cycloheteroalkyl, cyano. These groups may be attached to any carbon atom of the lower alkyl moiety.

The term "lower alkenyl" refers to a straight chain or branched hydrocarbon of two to ten carbon atoms having at least one carbon to carbon double bond.

The term "substituted lower alkenyl" refers to lower alkenyl as just described including one or more groups such as hydroxyl, thiol, alkylthiol, halogen, alkoxy, amino, amido, carboxyl, cycloalkyl, substituted cycloalkyl, heterocycle, cycloheteroalkyl, substituted cycloheteroalkyl, acyl, carboxyl, aryl, substituted aryl, aryloxy, hetaryl, substituted hetaryl, aralkyl, heteroaralkyl, alkyl, alkenyl, alkynyl, alkyl alkenyl, alkyl alkynyl, alkyl cycloalkyl, alkyl cycloheteroalkyl, cyano. These groups may be attached to any carbon atom to produce a stable compound.

The term "lower alkynyl" refers to a straight chain or branched hydrocarbon of two to ten carbon atoms having at least one carbon to carbon triple bond.

The term "substituted lower alkynyl" refers to lower alkynyl as just described including one or more groups such as hydroxyl, thiol, alkylthiol, halogen, alkoxy, amino, amido, carboxyl, cycloalkyl, substituted cycloalkyl, heterocycle, cycloheteroalkyl, substituted cycloheteroalkyl, acyl, carboxyl, aryl, substituted aryl, aryloxy, hetaryl, substituted hetaryl, aralkyl, heteroaralkyl, alkyl, alkenyl, alkynyl, alkyl alkenyl, alkyl alkynyl, alkyl cycloalkyl, alkyl cycloheteroalkyl, cyano. These groups may be attached to any carbon atom to produce a stable compound.

The term "alkoxy" refers to the group -OR, where R is lower alkyl, substituted lower alkyl, acyl, aryl, substituted aryl, aralkyl, substituted aralkyl, heteroalkyl, heteroarylalkyl, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, or substituted cycloheteroalkyl as defined below.

The term "alkylthio" denotes the group -SR, -S(O)_{n=1-2} -R, where R is lower alkyl, substituted lower alkyl, aryl, substituted aryl, aralkyl or substituted aralkyl as defined below.

The term "acyl" refers to groups -C(O)R, where R is hydrogen, lower alkyl substituted lower alkyl, aryl, substituted aryl.

The term "aryloxy" refers to groups -OAr, where Ar is an aryl, substituted aryl, heteroaryl, or substituted heteroaryl group as defined below.

The term "amino" refers to the group NRR', where R and R' may independently be hydrogen, lower alkyl, substituted lower alkyl, aryl, substituted aryl, hetaryl, cycloalkyl, or substituted hetaryl as defined below or acyl.

The term “amido” refers to the group -C(O)NRR', where R and R' may independently be hydrogen, lower alkyl, substituted lower alkyl, aryl, substituted aryl, hetaryl, substituted hetaryl as defined below.

5 The term “carboxyl” refers to the group -C(O)OR, where R may independently be hydrogen, lower alkyl, substituted lower alkyl, aryl, substituted aryl, hetaryl, substituted hetaryl and the like as defined.

The terms “aryl” or “Ar” refer to an aromatic carbocyclic group having at least one aromatic ring (e.g., phenyl or biphenyl) or multiple condensed rings in which at least one ring is aromatic, (e.g., 1,2,3,4-tetrahydronaphthyl, naphthyl, anthryl, or phenanthryl, 9-fluorenyl etc.).

10 The term “substituted aryl” refers to aryl optionally substituted with one or more functional groups, e.g., halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, trifluoromethyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, substituted heterocycle, heteroaryl, substituted heteroaryl, heteroalkyl, substituted
15 heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, sulfamido or cyano.

The term “heterocycle” refers to a saturated, unsaturated, or aromatic carbocyclic group having a single ring (e.g., morpholino, pyridyl or furyl) or multiple condensed rings (e.g., naphthpyridyl, quinoxalyl, quinoliny, indoliziny, indanyl or benzo[b]thienyl) and having at least one hetero
20 atom, such as N, O or S, within the ring.

The term “substituted heterocycle” refers to heterocycle optionally substituted with, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, trifluoromethyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, substituted
25 heterocycle, heteroaryl, substituted heteroaryl, heteroalkyl, substituted heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, sulfamido or cyano and the like.

The terms "heteroaryl" or "hetaryl" refer to a heterocycle in which at least one heterocyclic ring is aromatic.

- 5 The term "substituted heteroaryl" refers to a heterocycle optionally mono or poly substituted with one or more functional groups, e.g., halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, trifluoromethyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, substituted heterocycle, heteroaryl, substituted heteroaryl, heteroalkyl, substituted heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, sulfamido or cyano and the like.
- 10 The term "aralkyl" refers to the group -R-Ar where Ar is an aryl group and R is lower alkyl or substituted lower alkyl group. Aryl groups can optionally be unsubstituted or substituted with, e.g., halogen, lower alkyl, alkoxy, alkyl thio, trifluoromethyl, amino, amido, carboxyl, hydroxyl, aryl, aryloxy, heterocycle, hetaryl, substituted hetaryl, nitro, cyano, alkylthio, thiol, sulfamido and the like.
- 15 The term "heteroalkyl" refers to the group -R-Het where Het is a heterocycle group and R is a lower alkyl group. Heteroalkyl groups can optionally be unsubstituted or substituted with e.g., halogen, lower alkyl, lower alkoxy, lower alkylthio, trifluoromethyl, amino, amido, carboxyl, hydroxyl, aryl, aryloxy, heterocycle, hetaryl, substituted hetaryl, nitro, cyano, alkylthio, thiol, sulfamido and the like.
- 20 The term "heteroarylalkyl" refers to the group -R-HetAr where HetAr is an heteroaryl group and R lower alkyl or substituted loweralkyl. Heteroarylalkyl groups can optionally be unsubstituted or substituted with, e.g., halogen, lower alkyl, substituted lower alkyl, alkoxy, alkylthio, aryl, aryloxy, heterocycle, hetaryl, substituted hetaryl, nitro, cyano, alkylthio, thiol, sulfamido and the like.
- 25 The term "cycloalkyl" refers to a cyclic or polycyclic alkyl group containing 3 to 15 carbon. For polycyclic groups, these may be multiple condensed rings in which one of the distal rings may be aromatic (e.g. tetrahydronaphthalene, etc.).

The term "substituted cycloalkyl" refers to a cycloalkyl group comprising one or more substituents with, e.g. halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, trifluoromethyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, sulfamido or cyano and the like.

The term "cycloheteroalkyl" refers to a cycloalkyl group wherein one or more of the ring carbon atoms is replaced with a heteroatom (e.g., N, O, S or P).

10 The term "substituted cycloheteroalkyl" refers to a cycloheteroalkyl group as herein defined which contains one or more substituents, such as halogen, lower alkyl, lower alkoxy, lower alkylthio, trifluoromethyl, amino, amido, carboxyl, hydroxyl, aryl, aryloxy, heterocycle, hetaryl, substituted hetaryl, nitro, cyano, alkylthio, thiol, sulfamido and the like.

15 The term "alkyl cycloalkyl" refers to the group -R-cycloalkyl where cycloalkyl is a cycloalkyl group and R is a lower alkyl or substituted lower alkyl. Cycloalkyl groups can optionally be unsubstituted or substituted with e.g. halogen, lower alkyl, lower alkoxy, lower alkylthio, trifluoromethyl, amino, amido, carboxyl, hydroxyl, aryl, aryloxy, heterocycle, hetaryl, substituted hetaryl, nitro, cyano, alkylthio, thiol, sulfamido and the like.

20 The terms "therapy" and "treatment," as used interchangeably herein, refer to an intervention performed with the intention of alleviating the symptoms associated with, preventing the development of, or altering the pathology of a disease, disorder or condition. Thus, the terms therapy and treatment are used in the broadest sense, and include the prevention (prophylaxis), moderation, reduction, and curing of a disease, disorder or condition at various stages. Those in need of therapy/treatment include those already having the disease, disorder or condition as well as those prone to, or at risk of developing, the disease, disorder or condition and those in whom the disease, disorder or condition is to be prevented.

25 The term "subject" or "patient," as used herein, refers to an animal in need of treatment.

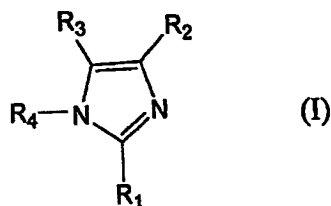
The term "animal," as used herein, refers to both human and non-human animals, including, but not limited to, mammals, birds and fish.

Administration of the compounds of the invention "in combination with" one or more further therapeutic agents, is intended to include simultaneous (concurrent) administration and
5 consecutive administration. Consecutive administration is intended to encompass various orders of administration of the therapeutic agent(s) and the compound(s) of the invention to the subject.

As used herein, the term "about" refers to a +/-10% variation from the nominal value. It is to be understood that such a variation is always included in any given value provided herein, whether or not it is specifically referred to.

10 **I. 2,4,5-Trisubstituted Imidazole Compounds**

The present invention provides compounds of the general formula (I):



15 or a salt thereof, wherein:

R1 is aryl, substituted aryl, heterocycle, substituted heterocycle, heteroaryl, or substituted heteroaryl;

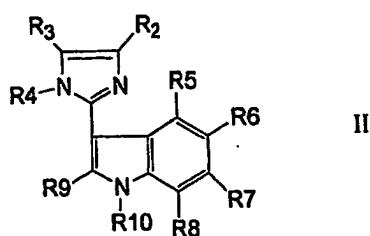
R2 and R3 are independently aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, or substituted heteroaryl or R2 and R3 when taken together along with the carbon
20 atoms they are attached to, form aryl or substituted aryl, heterocycle, substituted heterocycle, heteroaryl, or substituted heteroaryl;

R4 is hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle,

heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano.

In another embodiment of the present invention, the compound of formula (I) is other than Nortopsentin A, Nortopsentin B, Nortopsentin C and Nortopsentin D.

- 5 In another embodiment of the present invention, the compound of Formula I include the compound of the structural formula:

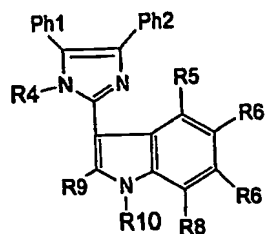


or a salt thereof, wherein:

- 10 R2 and R3 are independently aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, or substituted heteroaryl or R2 and R3 when taken together along with the carbon atoms they are attached form aryl or substituted aryl, heterocycle, heteroaryl, substituted heterocycle, or substituted heteroaryl;
- R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano;
- 15 20 R10 is H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, acyl, -CH₂-aryl, -CH₂-heteroaryl.

In another embodiment of the invention, the compound of Formula II is other than Nortopsentin A, Nortopsentin B, Nortopsentin C and Nortopsentin D.

- 25 In another embodiment of the present invention, the compound of Formula II include the compound of the structural formula III:

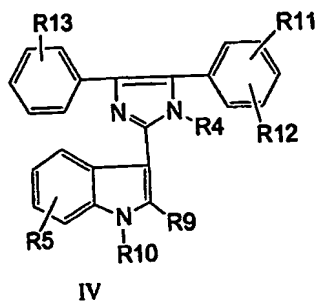


III

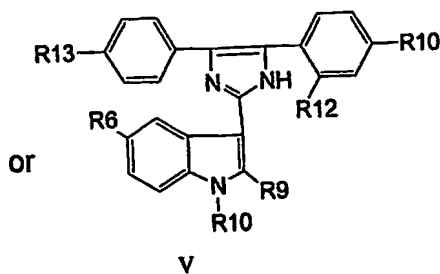
or a salt thereof, wherein:

- 5 Ph1 and Ph2 are independently selected from phenyl and substituted phenyl;
- R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle,
- 10 heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano;
- R10 is H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, acyl.

- 15 In another embodiment of the invention, the compound of Formula III is selected from:



IV



V

or a salt thereof, wherein:

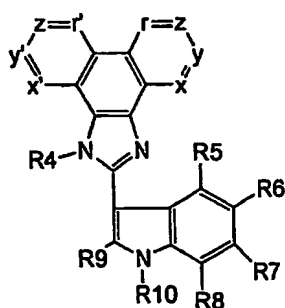
- R5, R6, R9, R11, R12 and R13 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy,
- 20

amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano;

R10 is H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl,

5 substituted aryl, heteroaryl, substituted heteroaryl, acyl.

In another embodiment of the present invention, the compound of Formula I include the compound of the structural formula:



VI

10 or a salt thereof, wherein:

R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano;

x is CR11 or N;

y is CR12 or N;

z is CR13 or N;

20 r is CR14 or N;

x' is CR15 or N;

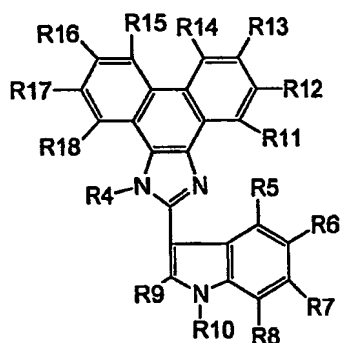
y' is CR16 or N;

z' is CR17 or N;

x' is CR18 or N;

R11, R12, R13, R14, R15, R16, R17 and R18 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano.

In another embodiment of the present invention, the compound of Formula I include the compound of the structural formula:



10

or a salt thereof, wherein:

R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano;

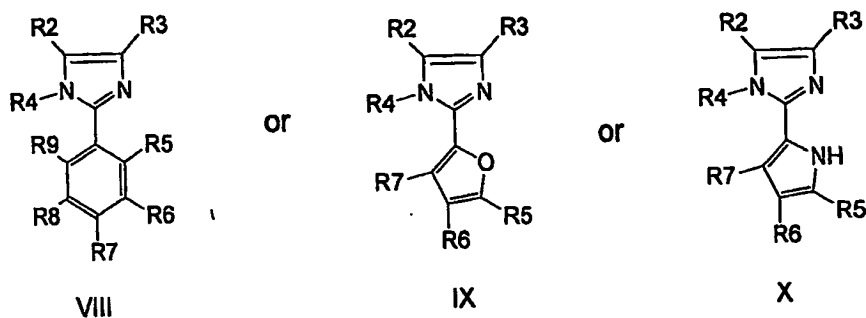
R10 is H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, acyl;

R11, R12, R13, R14, R15, R16, R17 and R18 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted

heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano.

In another embodiment of the present invention, the compound of Formula I is selected from the compound of formula:

5

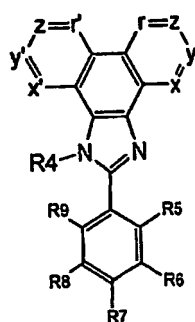


wherein:

R2 and R3 are independently aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, or substituted heteroaryl or R2 and R3 when taken together along with the carbon atoms they are attached to, form a aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, or substituted heteroaryl;

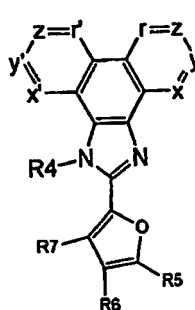
R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano.

In another embodiment of the present invention, the compound of Formula I include the compound of the structural formula:



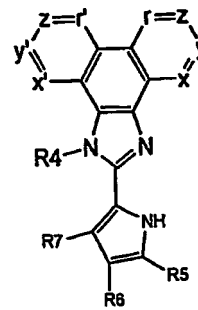
XI

or



XII

or



XIII

or a salt thereof, wherein:

R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol,
 5 lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl,
 substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino,
 amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle,
 heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or
 cyano;

10 x is CR11 or N;

y is CR12 or N;

z is CR13 or N;

r is CR14 or N;

x' is CR15 or N;

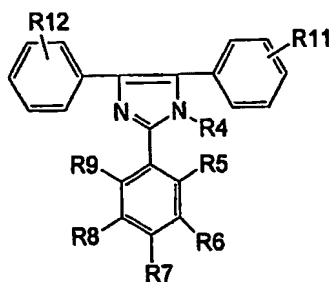
15 y' is CR16 or N;

z' is CR17 or N;

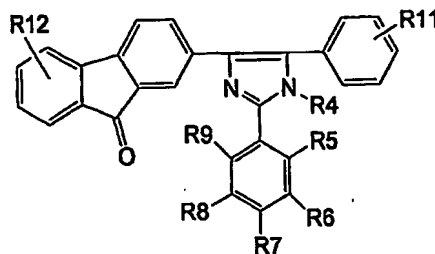
x' is CR18 or N;

R11, R12, R13, R14, R15, R16, R17 and R18 are independently selected from hydrogen,
 halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower
 20 alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio,
 acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted
 heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl,
 alkylcycloheteroalkyl, nitro, or cyano.

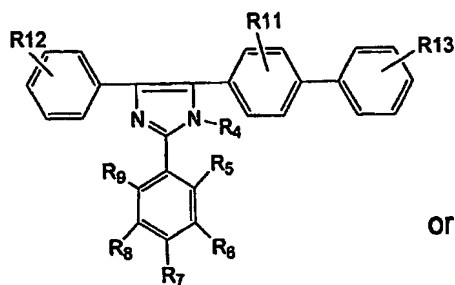
In another embodiment of the invention, the compound of Formula I is selected from:



XIV

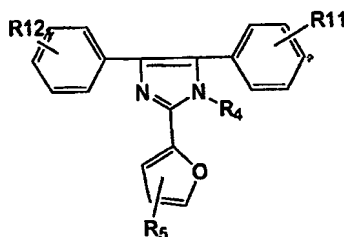


XV



XVI

OR



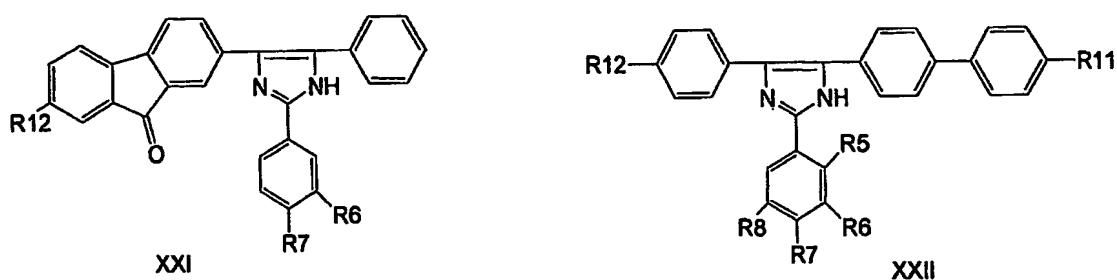
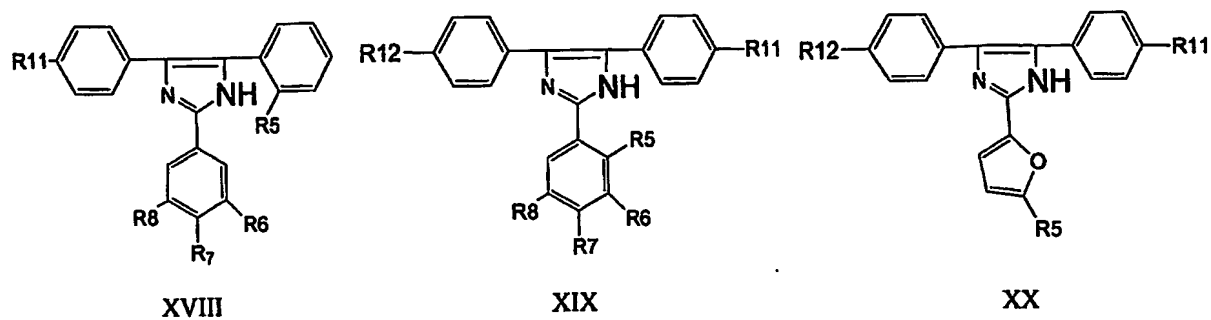
XVII

wherein:

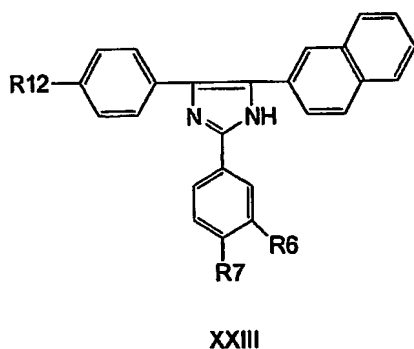
5 R4, R5, R6, R7, R8, R9, R11, R12 and R13 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, 10 aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano.

In another embodiment of the invention the compound of Formula I is selected from:

15



or



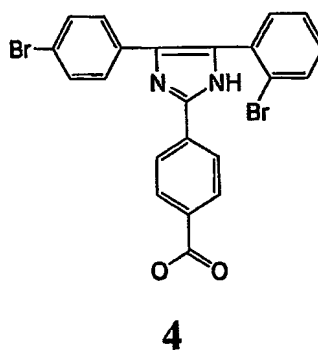
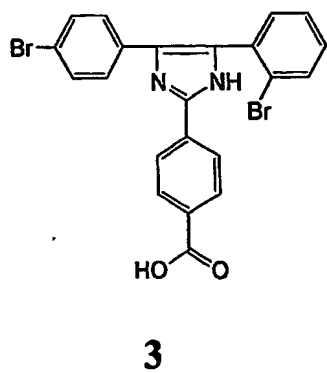
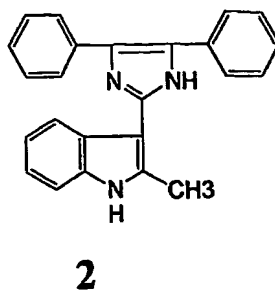
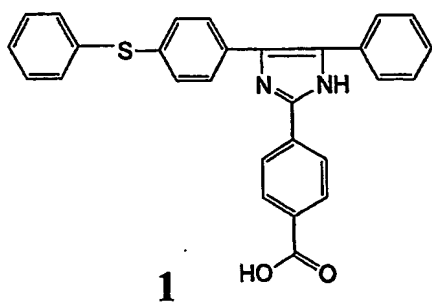
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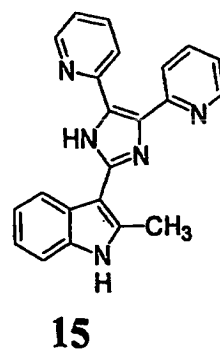
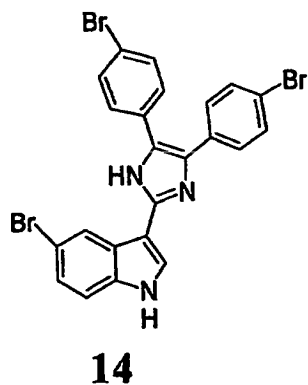
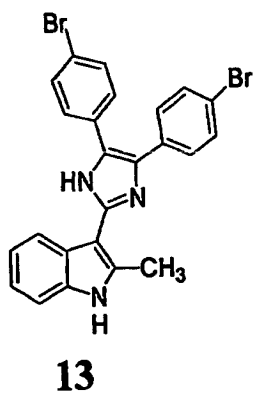
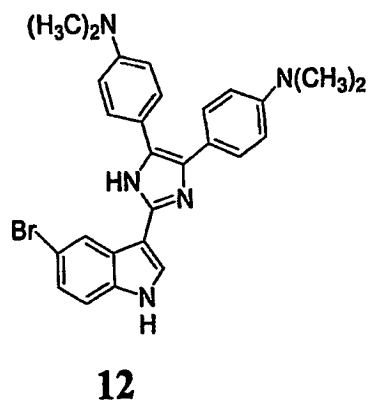
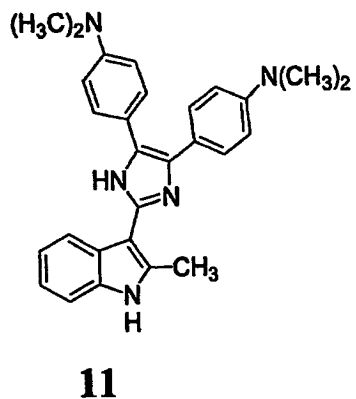
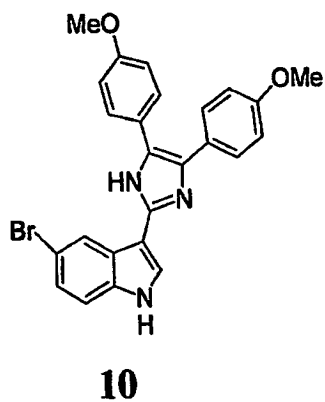
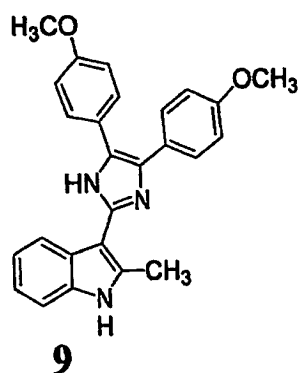
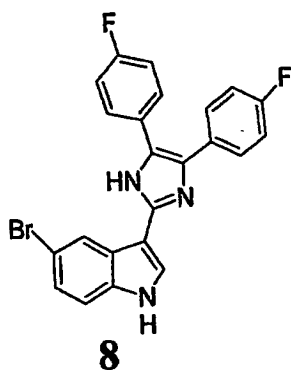
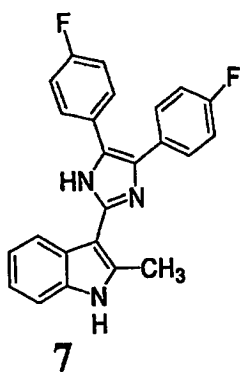
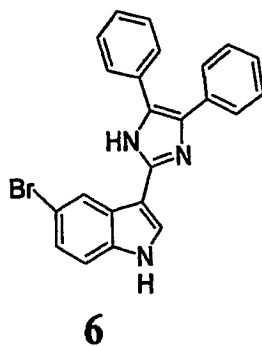
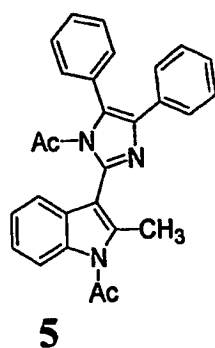
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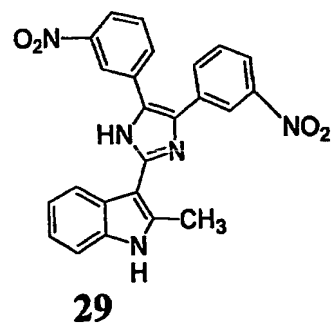
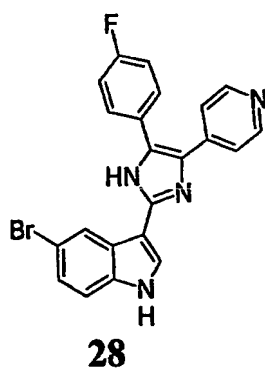
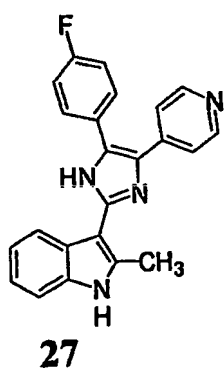
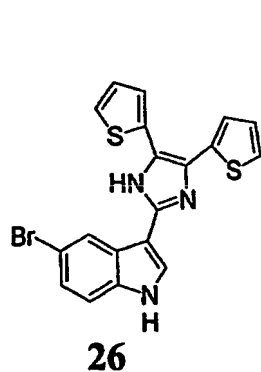
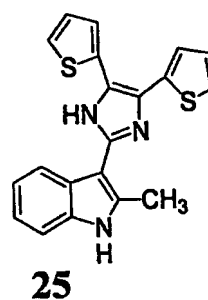
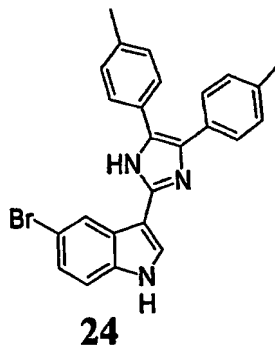
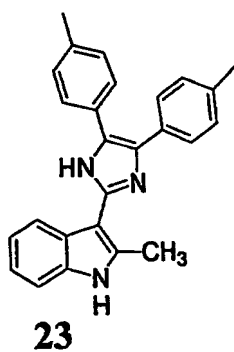
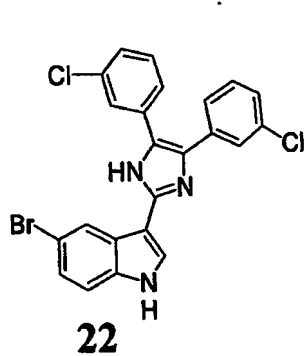
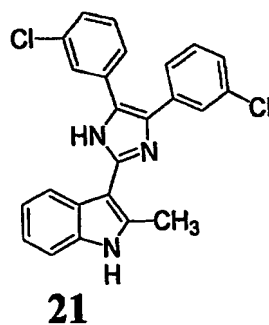
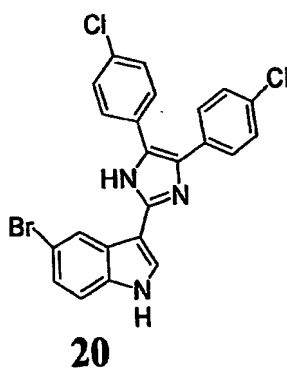
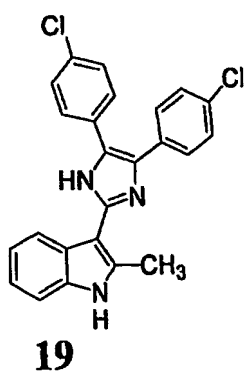
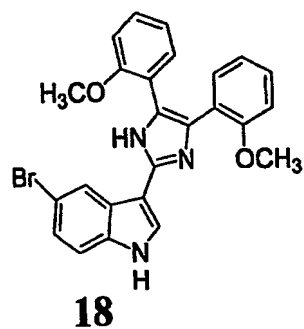
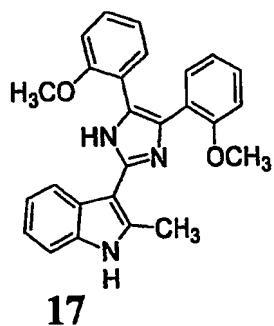
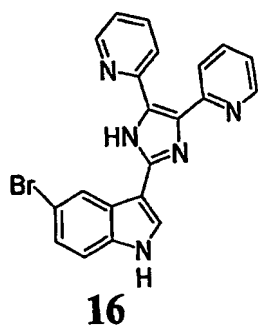
R5, R6, R7, R8, R9, R11 and R12 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano.

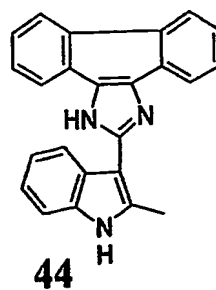
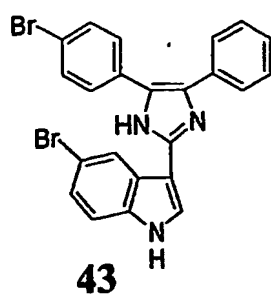
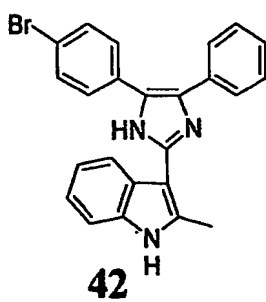
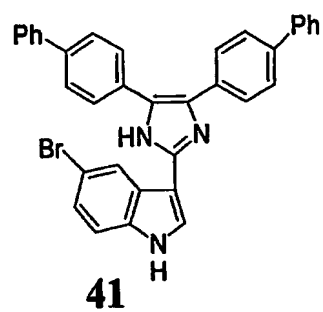
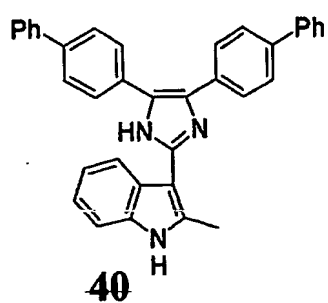
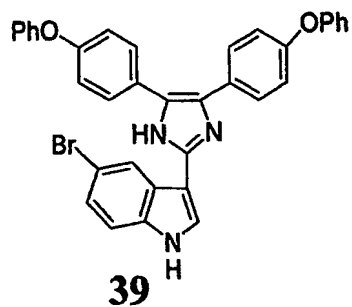
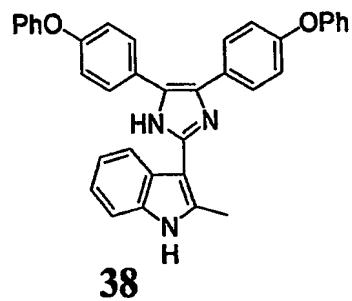
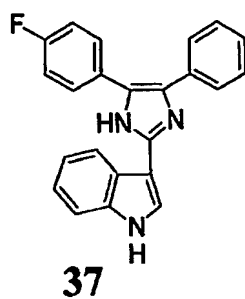
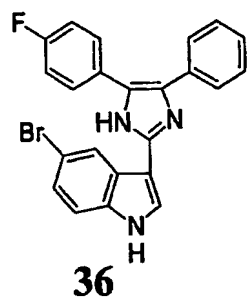
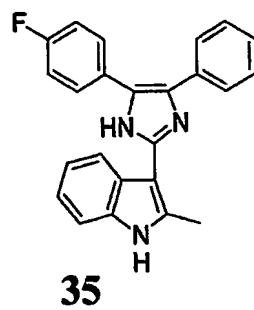
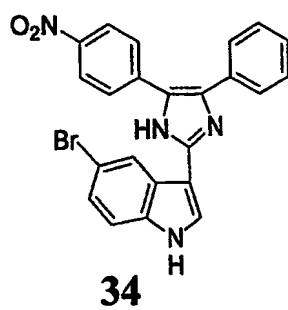
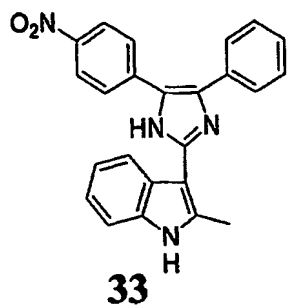
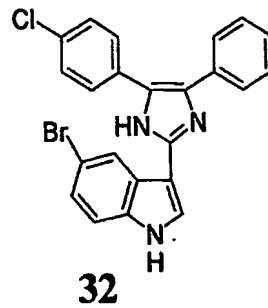
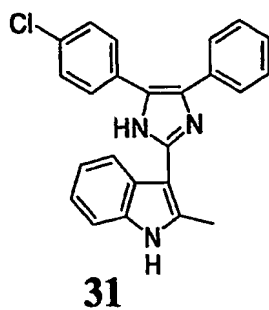
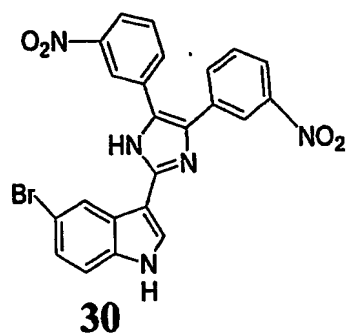
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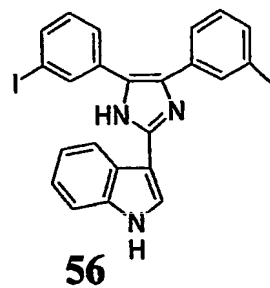
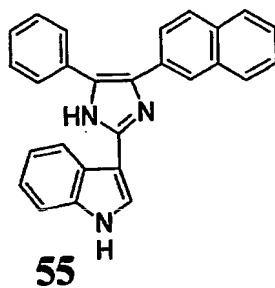
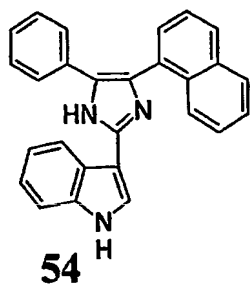
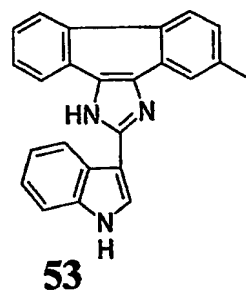
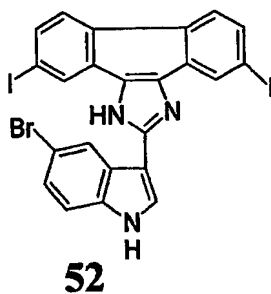
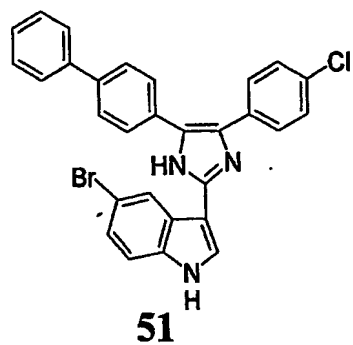
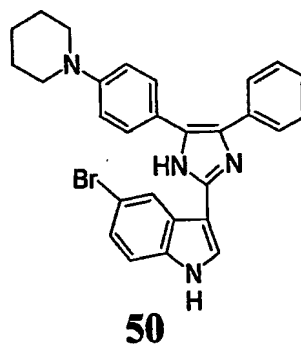
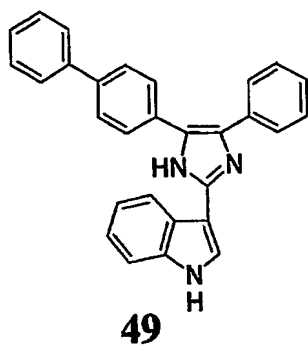
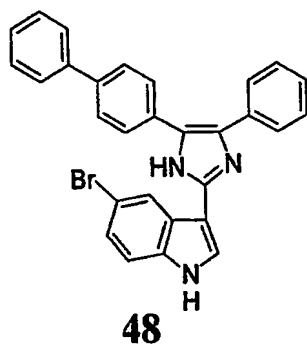
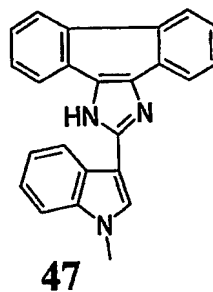
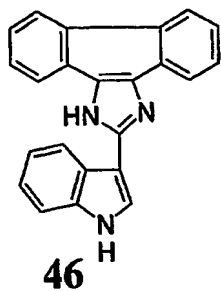
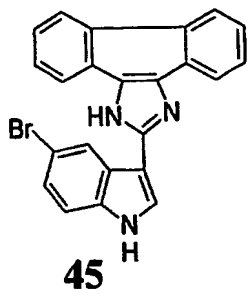
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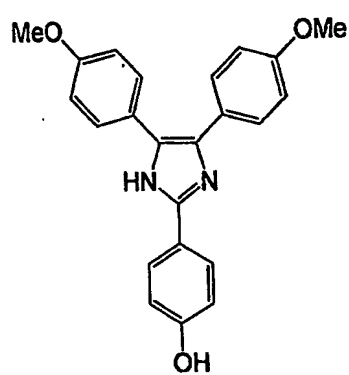




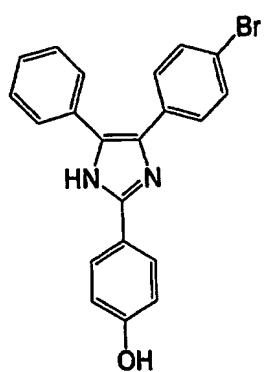




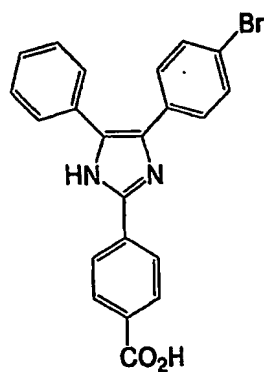




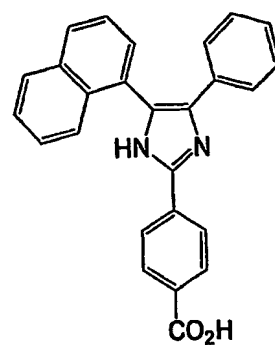
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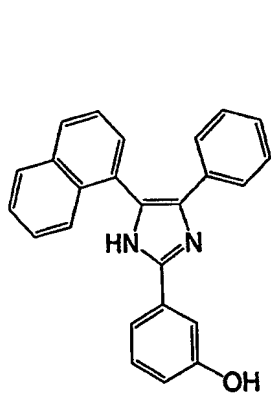
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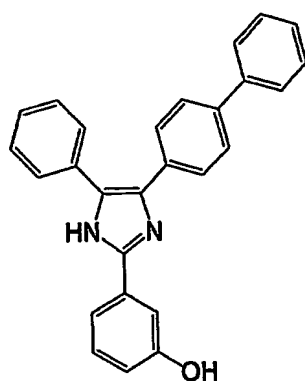
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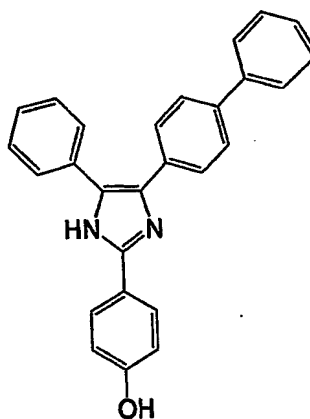
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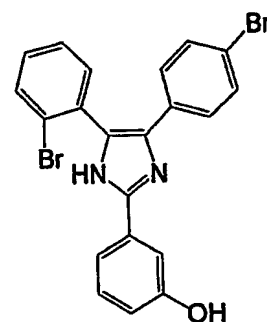
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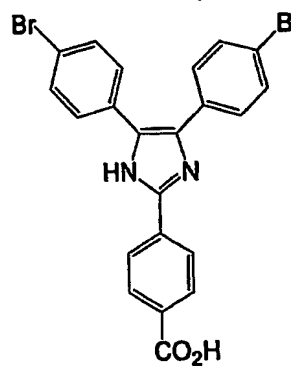
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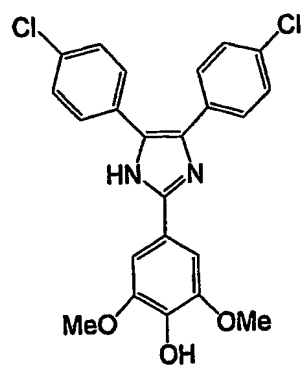
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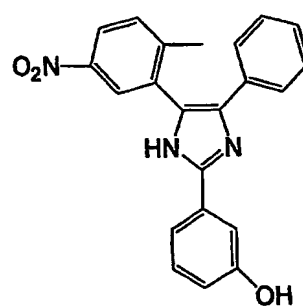
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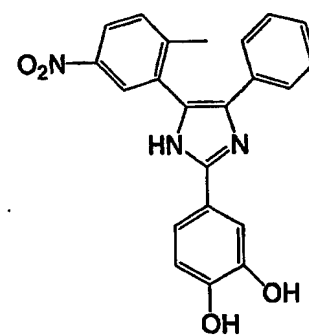
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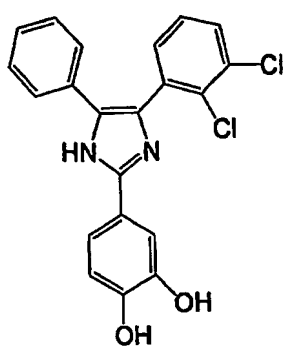
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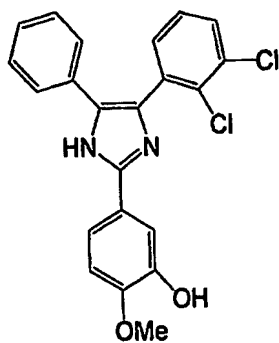
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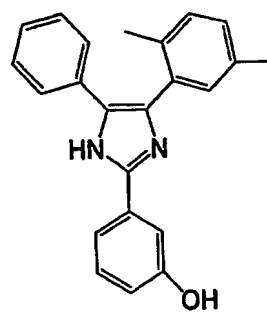
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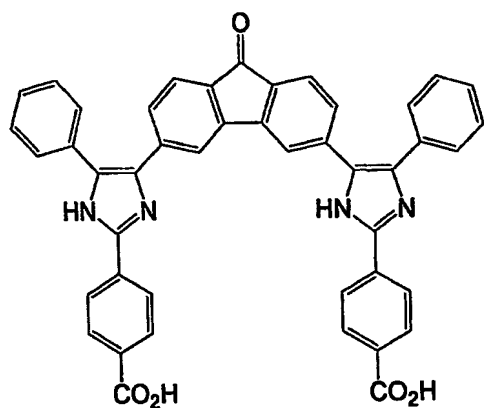
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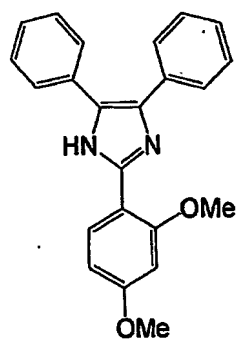
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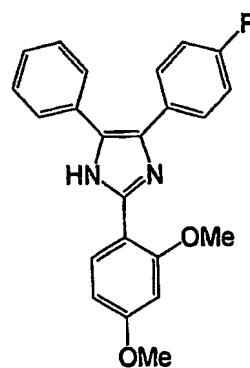
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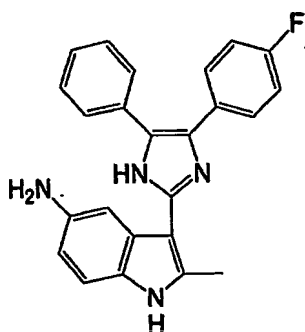
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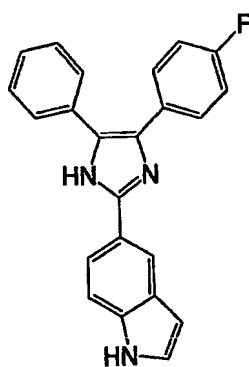
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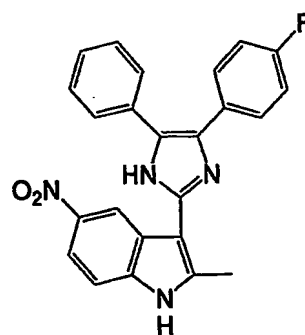
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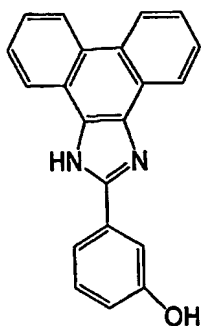
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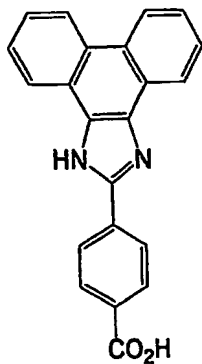
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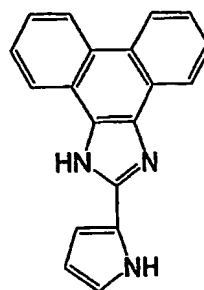
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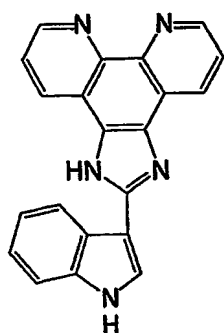
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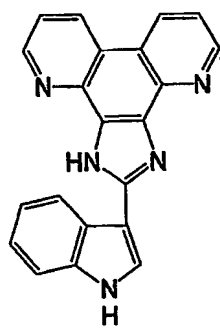
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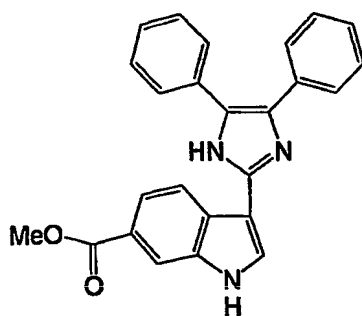
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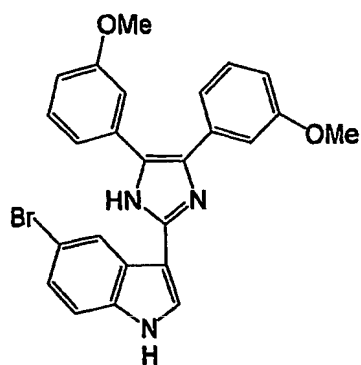
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The present invention includes pharmaceutically acceptable salts of the compounds defined by Formula I. Compounds according to the present invention can possess a sufficiently acidic, a

sufficiently basic, or both functional groups, and accordingly react with a number of organic and inorganic bases, and organic and inorganic acids, to form pharmaceutically acceptable salts.

The term "pharmaceutically acceptable salt" as used herein, refers to a salt of a compound of Formula I, which is substantially non-toxic to living organisms. Typical pharmaceutically acceptable salts include those salts prepared by reaction of the compound of the present invention with a pharmaceutically acceptable mineral or organic acid or an organic or inorganic base. Such salts are known as acid addition and base addition salts.

Acids commonly employed to form acid addition salts are inorganic acids such as hydrochloric acid, hydrobromic acid, hydroiodic acid, sulphuric acid, phosphoric acid, and the like, and organic acids such as *p*-toluenesulphonic acid, methanesulphonic acid, oxalic acid, *p*-bromophenylsulphonic acid, carbonic acid, succinic acid, citric acid, benzoic acid, acetic acid, and the like. Examples of such pharmaceutically acceptable salts are the sulphate, pyrosulphate, bisulphate, sulphite, phosphate, monohydrogenphosphate, dihydrogenphosphate, metaphosphate, pyrophosphate, bromide, iodide, acetate, propionate, decanoate, caprylate, acrylate, formate, hydrochloride, dihydrochloride, isobutyrate, caproate, heptanoate, propiolate, oxalate, malonate, succinate, suberate, sebacate, fumarate, maleate, butyne-1,4-dioate, hexyne-1,6-dioate, benzoate, chlorobenzoate, methylbenzoate, hydroxybenzoate, methoxybenzoate, phthalate, xylenesulphonate, phenylacetate, phenylpropionate, phenylbutyrate, citrate, lactate, gamma-hydroxybutyrate, glycolate, tartrate, methanesulphonate, propanesulphonate, naphthalene-1-sulfonate, naphthalene-2-sulfonate, mandelate and the like. Preferred pharmaceutically acceptable acid addition salts are those formed with mineral acids such as hydrochloric acid and hydrobromic acid, and those formed with organic acids such as maleic acid and methanesulphonic acid.

Salts of amine groups may also comprise quarternary ammonium salts in which the amino nitrogen carries a suitable organic group such as an alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, or aralkyl moiety.

Base addition salts include those derived from inorganic bases, such as ammonium or alkali or alkaline earth metal hydroxides, carbonates, bicarbonates, and the like. Bases useful in preparing the salts of this invention thus include sodium hydroxide, potassium hydroxide, ammonium

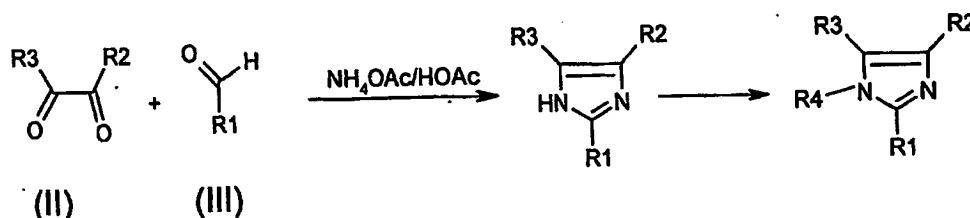
hydroxide, potassium carbonate, sodium carbonate, sodium bicarbonate, potassium bicarbonate, calcium hydroxide, calcium carbonate, and the like.

- One skilled in the art will understand that the particular counterion forming a part of a salt of this invention is usually not of a critical nature, so long as the salt as a whole is pharmacologically acceptable and as long as the counterion does not contribute undesired qualities to the salt as a whole. The present invention further encompasses the pharmaceutically acceptable solvates of a compound of Formula I. Many of the compounds of Formula I can combine with solvents such as water, methanol, ethanol and acetonitrile to form pharmaceutically acceptable solvates such as the corresponding hydrate, methanolate, ethanolate and acetonitrilate.
- 10 The compounds of the present invention may have multiple asymmetric (chiral) centres. As a consequence of these chiral centres, the compounds of the present invention occur as racemates, mixtures of enantiomers and as individual enantiomers, as well as diastereomers and mixtures of diastereomers. All asymmetric forms, individual isomers and combinations thereof, are within the scope of the present invention.
- 15 It will be readily understood by one skilled in the art that if the stereochemistry of a compound of Formula I is critical to its activity, then the relative stereochemistry of the compound is established early during synthesis to avoid subsequent stereoisomer separation problems. Further manipulation of the molecule will then employ stereospecific procedures so as to maintain the desired chirality.
- 20 Non-toxic metabolically-labile esters or amides of a compound of Formula I are those that are hydrolysed *in vivo* to afford the compound of Formula I and a pharmaceutically acceptable alcohol or amine. Examples of metabolically-labile esters include esters formed with (1-6C) alkanols, in which the alkanol moiety may be optionally substituted by a (1-8C) alkoxy group, for example methanol, ethanol, propanol and methoxyethanol. Non-limiting examples of
- 25 metabolically-labile amides include amides formed with amines such as methylamine.

II. Preparation of Compounds of Formula I

As is known in the art, triaryl imidazole compounds can be prepared by a number of standard techniques. Compounds of Formula I, therefore, can be prepared by several general synthetic methods, for example, as described by Grimmett, (Grimmett, M.R., *Comprehensive Heterocyclic Chemistry: The Structure, Reaction, Synthesis and Uses of Heterocyclic Compounds*, A. R. Katrizky and C. W. Rees, eds., Vol. 5, Pergamon Press. Oxford, 1984, pp. 457-498; Grimmett, M. R., *Imidazole and Benzimidazole Synthesis*, Academic Press, San Diego CA, 1997).

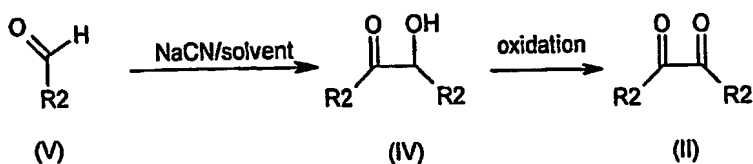
In one embodiment of the present invention, compounds of Formula I are prepared *via* solution or solid phase synthesis, by reacting a dione of Formula II with the aldehyde (III) at elevated temperature in the presence of ammonium acetate in acetic acid (see, for example, Krieg *et al.*, *Naturforsch.* 1967, **22b**:132; Sarshar *et al.*, *Tetrahedron Lett.* 1996, **37**:835-838).



The compounds of Formula (II) and (III) are either commercially available or may be prepared using standard procedures known to a person skilled in the relevant art. Compounds of Formula II, therefore, can be prepared by several general synthetic methods, for example, as described by: Fischer *et. al* (*J. Am. Chem. Soc.* 1961, **83**, 4208-4210); Guijarro *et al.* (*J. Am. Chem. Soc.* 1999, **121**, 4155-4157); Chi *et. al.* (*Synth. Comm.* 1994, **24**(15), 2119-2122) and Armesto *et. al.* (*Synthesis*, 1988, 799-801).

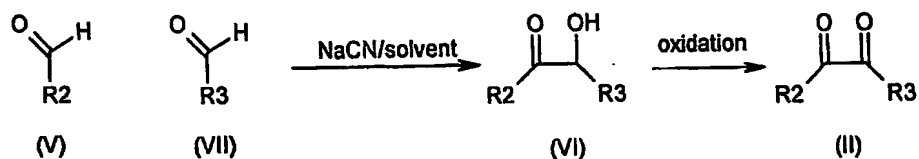
Compounds of formula II can also be prepared:

- i) by oxidizing a compound of formula (IV). Compounds of formula (IV), in turn can be prepared by reacting a compounds of formula (V) with sodium cyanide in the presence of a solvent as shown below, wherein R3 = R2 and R2 is as defined above:



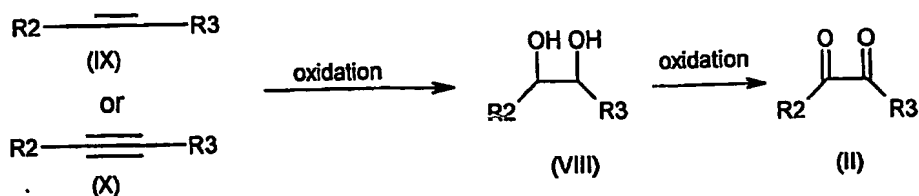
or,

- 5 ii) by oxidizing a compound of formula (VI). Compounds of formula (VI), in turn can be prepared by treating a compound of formula (V) and a compound of formula (VII) with sodium cyanide in the presence of a solvent as shown below, wherein R₂ and R₃ are as defined above:



or,

- 15 iii) by oxidizing a compound of formula (VIII). Compounds of formula (VIII) in turn can be prepared by oxidizing a compound of formula (IX) or (X) as shown below, wherein R₂ and R₃ are as defined above:

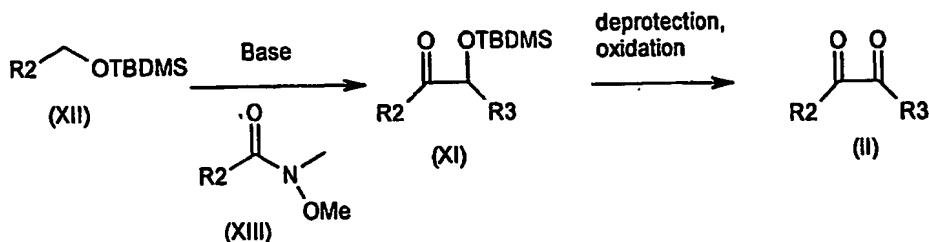


or,

- iv) by oxidizing a compound of formula (X) using PdCl₂ in DMSO,

20 or,

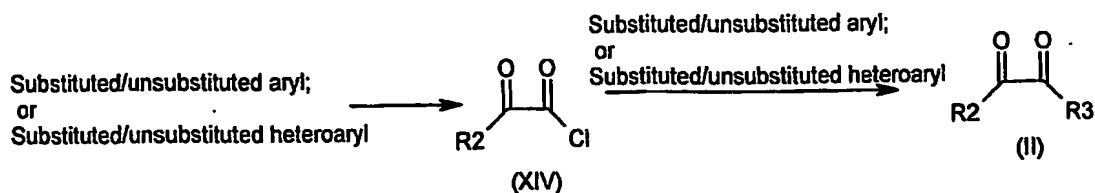
- v) by deprotecting and oxidizing a compound of formula (XI). Compounds of formula (XI) in turn can be prepared by reacting a compound of formula (XII) with a compound of formula (XIII) in the presence of a suitable base:



wherein R^2 and R^3 are independently aryl, substituted aryl, heteroaryl or substituted heteroaryl,

or,

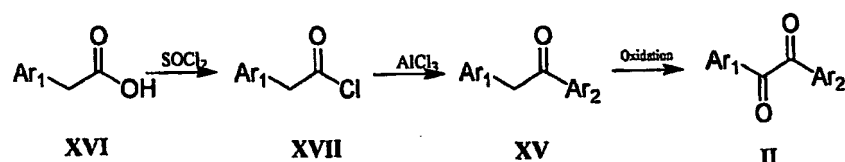
- vi) by reacting a compound of formula (XIV) with a substituted or unsubstituted aryl or substituted or unsubstituted heteroaryl under Friedel-Crafts acylation conditions or by nucleophilic displacement of the chloride in compound of formula XIV. Compounds of formula (XIV) in turn can be prepared by reacting a substituted or unsubstituted aryl or substituted or unsubstituted heteroaryl with oxalyl chloride under Friedel-Crafts acylation conditions:



wherein R^2 and R^3 are independently aryl, substituted aryl, heteroaryl or substituted heteroaryl;

or

- vii) by oxidising a compound of formula (XV). Compounds of formula (XV) in turn are prepared by reacting a compound of formula (XVI) with thionyl chloride in benzene with catalytic dimethylformamide to form an intermediate (XVII). This intermediate (XVII) is then used directly without purification in a Freidel–Crafts reaction to produce the ketone (XV).



10 III. Anti-cancer Activity of Compounds of Formula I

The ability of a candidate compound of Formula I to inhibit neoplastic cell growth and/or proliferation can be tested using standard techniques known in the art. In addition, compounds of Formula I that demonstrate inhibitory activity may be further tested *in vitro* and/or *in vivo* in combination with various known chemotherapeutics to evaluate their potential use in combination therapies. Exemplary methods of testing candidate compounds of Formula I are provided below and in the Examples included herein. One skilled in the art will understand that other methods of testing the compounds are known in the art and are also suitable for testing candidate compounds.

A. In vitro Testing

- 20 Candidate compounds of Formula I can be assayed initially *in vitro* for their ability to inhibit cell growth (*i.e.* their cytotoxicity) using standard techniques. In general, cells of a specific test cell line (typically a cancer cell line) are grown to a suitable density (*e.g.* approximately 1×10^4) and the candidate compound is added. After an appropriate incubation time (typically between about 48 to 74 hours), cell survival is assessed, for example, by assaying for tetrazolium salt (or modified tetrazolium salt) cleavage, or by using the resazurin reduction test (see Fields & Lancaster (1993) *Am. Biotechnol. Lab.* 11:48–50; O'Brien *et al.*, (2000) *Eur. J. Biochem.* 267:5421–5426 and U.S. Patent No. 5,501,959), the sulforhodamine assay (Rubinstein *et al.*, (1990) *J. Natl. Cancer Inst.* 82:113–118) or the neutral red dye test (Kitano *et al.*, (1991) *Euro.*

J. Clin. Investg. 21:53-58; West *et al.*, (1992) *J. Investigative Derm.* 99:95-100). Inhibition of cell growth is determined by comparison of cell survival in the treated culture with cell survival in one or more control cultures, for example, cultures not pre-treated with the candidate compound and/or those pre-treated with a control compound (typically a known therapeutic).

5 Other suitable techniques for assessing cytotoxicity are known in the art.

Assays that measure metabolic activity (such as tetrazolium-based assays) can also be used to assess the effect of candidate compounds on cell activation and /or proliferation, due the fact that proliferating cells are metabolically more active than resting cells.

10 Candidate compounds can also be tested *in vitro* for their ability to inhibit anchorage-independent growth of tumour cells. Anchorage-independent growth is known in the art to be a good indicator of tumourigenicity. In general, anchorage-independent growth is assessed by plating cells from an appropriate cancer cell-line onto soft agar and determining the number of colonies formed after an appropriate incubation period. Growth of cells treated with the candidate compound can then be compared with that of cells treated with an appropriate control
15 (as described above).

A variety of cancer cell-lines suitable for testing the candidate compounds are known in the art. In one embodiment of the present invention, *in vitro* testing of the candidate compounds is conducted in a human cancer cell-line. Examples of suitable human cancer cell-lines for *in vitro* testing of the compounds of the present invention include, but are not limited to, colon and
20 colorectal carcinoma cell lines such as HT-29, CaCo, LoVo, COLO320 and HCT-116; non small cell lung cancer cell lines such as NCI-H460, small cell lung cancer cell lines such as H209; breast cancer cell lines such as MCF-7, T47D and MDA-MB-231; ovarian cancer cell lines such as SK-OV-3; prostate cancer cell lines such as PC-3 and DU-145; chronic myeloid leukaemia cell lines such as K562; bladder cancer cell lines such as T24; brain cancer cell lines such as U-
25 87-MG; pancreatic cancer cell lines such as AsPC-1, SU.86.86 and BxPC-3; kidney cancer cell lines such as A498 and Caki-1; liver cancer cell lines such as HepG2, and skin cancer cell lines such as A2058 and C8161.

As is well known in the art, a compound that is to become a viable anti-cancer drug should demonstrate some level of selective action toward neoplastic (or cancer) cells in comparison to

normal proliferating cells. It may also be desirable, therefore, to test the selectivity of the candidate compounds of Formula I. An exemplary method of assessing the differential sensitivity between normal and cancer cells for a compound has been described by Vassilev *et al.* (*Anti-Cancer Drug Design* (2001) 16:7). This method involves the comparison of IC₉₀ values, *i.e.* the molar concentration of a test compound required to cause 90% growth inhibition of exponentially growing cells. The IC₉₀ value is believed to more accurately reflect the ability of a compound to induce apoptosis than other measurements. Thus, the IC₉₀ values for candidate compounds can be evaluated in various cancer cell lines (such as those outlined above) and normal cells (such as HUVEC and/or WI38 cells) and compared. IC₉₀ values can be measured using a variety of standard techniques including those described above for cytotoxicity testing. In one embodiment of the present invention, candidate compounds demonstrating an IC₉₀ value about 2-fold greater for one or more cancer cell line when compared to a normal control cell line are selected for further testing.

While the mechanism of action of the compounds of Formula I is not relevant to the instant invention, assays to investigate potential mechanisms of action of the compounds may be conducted if desired in order to provide information useful in determining what aspects of tumour growth the compounds affect. This type of information may help to determine cancer types that will benefit from treatment with the compounds. Examples of such assays include, but are not limited to, cell-cycle analysis (for example, employing flow cytometry techniques), apoptosis assays (such as DNA fragmentation analysis), anti-angiogenesis assays (for example, various Matrigel assays, including cord formation and Matrigel plug assays) and immunohistochemical analysis.

Toxicity of the candidate compounds can also be initially assessed *in vitro* using standard techniques. For example, human primary fibroblasts can be treated *in vitro* with a compound of Formula I and then tested at different time points following treatment for their viability using a standard viability assay, such as the assays described above or the trypan-blue exclusion assay.

- 5 Cells can also be assayed for their ability to synthesize DNA, for example, using a thymidine incorporation assay, and for changes in cell cycle dynamics, for example, using a standard cell sorting assay in conjunction with a fluorocytometer cell sorter (FACS).

B. In vivo Testing

The ability of the candidate compounds to inhibit tumour growth, proliferation and/or metastasis *in vivo* can be determined in an appropriate animal model using standard techniques known in the art (see, for example, Enna, *et al.*, *Current Protocols in Pharmacology*, J. Wiley & Sons, Inc., New York, NY). Non-limiting examples are provided in Table 1.

In general, current animal models for screening anti-tumour compounds are xenograft models, in which a human tumour has been implanted into an animal. For example, the candidate compounds can be tested *in vivo* on solid tumours using mice that are subcutaneously grafted or injected with 30 to 60 mg of a tumour fragment, or an appropriate number of tumour cells (*e.g.* about 10^6 to 10^7) on day 0. The animals bearing tumours are mixed before being subjected to the various treatments and controls. In the case of treatment of advanced tumours, tumours are allowed to develop to the desired size, animals having insufficiently developed tumours being eliminated. The selected animals are distributed at random to undergo the treatments and controls. Animals not bearing tumours may also be subjected to the same treatments as the tumour-bearing animals in order to be able to dissociate the toxic effect from the specific effect on the tumour. Chemotherapy generally begins from 3 to 22 days after grafting, depending on the type of tumour, and the animals are observed every day. Candidate compounds can be administered to the animals, for example, by bolus infusion. The different animal groups are weighed about 3 or 4 times a week until the maximum weight loss is attained, after which the groups are weighed at least once a week until the end of the trial.

The tumours are measured about 2 or 3 times a week until the tumour reaches a pre-determined size and / or weight, or until a pre-determined time period has passed, or until the animal dies (if this occurs before the tumour reaches the pre-determined size / weight). The animals are then sacrificed and the tissue histology, size and / or proliferation of the tumour assessed.

If desired, one or more standard immunohistochemical tests may also be conducted on tissues isolated from the test animals in order to determine the effects of the compound on tumour growth, differentiation, apoptosis and/or angiogenesis. Examples of such tests include, but are not limited to, the use of specific antibodies (for example, antibodies against Ki-67 to assess proliferation, CD31 to assess angiogenesis, NK1.1 as an indication of the presence of NK cells,

F4/80 as an indication of the presence of macrophages) and TUNEL assays to determine apoptosis.

5 Other models, such as orthopedic implantation of tumours into animals (*i.e.* the implantation of cancer cells of a certain type into the corresponding tissue in the animal, such as pancreatic cancer cells into the pancreas), may also be used to assess the effect of the candidate compounds on tumour growth and proliferation. In addition, the effect of the candidate compound on spontaneous tumours in normal mice can be assessed.

10 For the study of the effect of the candidate compounds on lymphomas or leukaemias, the animals are grafted or injected with a particular number of cells, and the anti-tumour activity is determined by the increase in the survival time of the treated mice relative to the controls.

To study the effect of the candidate compounds on tumour metastasis, tumour cells are typically treated with the compound *ex vivo* and then injected into a suitable test animal. The spread of the tumour cells from the site of injection is then monitored over a suitable period of time.

15 *In vivo* toxic effects of the compounds of Formula I can be evaluated by standard techniques, for example, by measuring their effect on animal body weight during treatment and by performing haematological profiles and liver enzyme analysis after the animal has been sacrificed (survival assays).

Table 1: Examples of *in vivo* models of human cancer

| Cancer Model | Cell Type |
|---|-------------------------------------|
| Tumour Growth Assay | Prostate (PC-3, DU145) |
| Human solid tumour xenografts in mice (sub-cutaneous injection) | Breast (MDA-MB-231, MVB-9) |
| | Colon (HT-29) |
| | Lung (NCI-H460, NCI-H209) |
| | Pancreatic (ASPC-1, SU86.86) |
| | Pancreatic: drug resistant (BxPC-3) |
| | Skin (A2058, C8161) |
| | Cervical (SIHA, HeLa-S3) |
| | Cervical: drug resistant (HeLa S3- |

| Cancer Model | Cell Type |
|--|---|
| | HU-resistance) Liver (HepG2) Brain (U87-MG) . Renal (Caki-1, A498) Ovary (SK-OV-3) Bladder (T24) |
| Tumour Growth Assay Human solid tumour isografts in mice (fat pad injection) | Breast: drug resistant (MDA-CDDP-S4, MDA-MB435-To.1) |
| Survival Assay Experimental model of lymphoma and leukaemia in mice | Human: Burkitts lymphoma (Non-Hodgkin's) (raji) Murine: erythroleukemia (CB7 Friend retrovirus-induced) |
| Experimental model of lung metastasis in mice | Human: melanoma (C8161) Murine: fibrosarcoma (R3) |

IV. Toxicity Testing

It is important that the anti-cancer compounds of the present invention exhibit low toxicity *in vivo*. Toxicity tests for potential drugs are well-known in the art (see, for example, Hayes, A.W., ed., (1994), *Principles and Methods of Toxicology*, 3rd ed., Raven Press, NY; Maines, M., ed., *Current Protocols in Toxicology*, John Wiley & Sons, Inc., NY).

In vitro acute toxicity testing of a compound of Formula I can be performed using mammalian cell lines (see, for example, Ekwall, B., *Ann. N.Y. Acad. Sci.*, (1983) 407:64-77). Selection of an appropriate cell line is dependent on the potential application of the candidate compound and can be readily determined by one skilled in the art.

In vivo toxicity testing can be performed by standard methodology. For example, by injecting varying concentrations of the candidate compound into an appropriate animal model. The compound can be injected once, or administration can be repeated over several days. The toxic effects of the compound can be evaluated over an appropriate time period by monitoring the

general health and body weight of the animals. After the completion of the period of assessment, the animals can be sacrificed and the appearance and weight of the relevant organs determined.

V. Therapeutic Uses of Compounds of Formula I

5 The compounds of Formula I can be used in the treatment and/or stabilisation of various types of cancers. The compounds can be used alone or they can be used as part of a multi-drug regimen in combination with one or more known therapeutics.

10 Examples of cancers which may be may be treated or stabilized in accordance with the present invention include, but are not limited to leukaemia, carcinomas, adenocarcinomas, melanomas and sarcomas. Carcinomas, adenocarcinomas and sarcomas are also frequently referred to as "solid tumours," examples of commonly occurring solid tumours include, but are not limited to, cancer of the brain, breast, cervix, colon, head and neck, kidney, lung, ovary, pancreas, prostate, stomach and uterus, non-small cell lung cancer and colorectal cancer.

15 The term "leukaemia" refers broadly to progressive, malignant diseases of the blood-forming organs. Leukaemia is typically characterized by a distorted proliferation and development of leukocytes and their precursors in the blood and bone marrow but can also refer to malignant diseases of other blood cells such as erythroleukaemia, which affects immature red blood cells. Leukaemia is generally clinically classified on the basis of (1) the duration and character of the disease – acute or chronic; (2) the type of cell involved – myeloid (myelogenous), lymphoid (lymphogenous) or monocytic, and (3) the increase or non-increase in the number of abnormal
20 cells in the blood – leukaemic or aleukaemic (subleukaemic). Leukaemia includes, for example, acute nonlymphocytic leukaemia, chronic lymphocytic leukaemia, acute granulocytic leukaemia, chronic granulocytic leukaemia, acute promyelocytic leukaemia, adult T-cell leukaemia, aleukaemic leukaemia, aleukocythemmic leukaemia, basophylic leukaemia, blast cell leukaemia, bovine leukaemia, chronic myelocytic leukaemia, leukaemia cutis, embryonal leukaemia, eosinophilic leukaemia, Gross' leukaemia, hairy-cell leukaemia, hemoblastic leukaemia,
25 hemocytoblastic leukaemia, histiocytic leukaemia, stem cell leukaemia, acute monocytic leukaemia, leukopenic leukaemia, lymphatic leukaemia, lymphoblastic leukaemia, lymphocytic leukaemia, lymphogenous leukaemia, lymphoid leukaemia, lymphosarcoma cell leukaemia, mast cell leukaemia, megakaryocytic leukaemia, micromyeloblastic leukaemia, monocytic leukaemia,

myeloblastic leukaemia, myelocytic leukaemia, myeloid granulocytic leukaemia, myelomonocytic leukaemia, Naegeli leukaemia, plasma cell leukaemia, plasmacytic leukaemia, promyelocytic leukaemia, Rieder cell leukaemia, Schilling's leukaemia, stem cell leukaemia, subleukaemic leukaemia, and undifferentiated cell leukaemia.

- 5 The term "sarcoma" generally refers to a tumour which originates in connective tissue, such as muscle, bone, cartilage or fat, and is made up of a substance like embryonic connective tissue and is generally composed of closely packed cells embedded in a fibrillar or homogeneous substance. Sarcomas include soft tissue sarcomas, chondrosarcoma, fibrosarcoma, lymphosarcoma, melanosarcoma, myxosarcoma, osteosarcoma, Abemethy's sarcoma, adipose
- 10 sarcoma, liposarcoma, alveolar soft part sarcoma, ameloblastic sarcoma, botryoid sarcoma, chloroma sarcoma, chorio carcinoma, embryonal sarcoma, Wilms' tumour sarcoma, endometrial sarcoma, stromal sarcoma, Ewing's sarcoma, fascial sarcoma, fibroblastic sarcoma, giant cell sarcoma, granulocytic sarcoma, Hodgkin's sarcoma, idiopathic multiple pigmented haemorrhagic sarcoma, immunoblastic sarcoma of B cells, lymphoma, immunoblastic sarcoma
- 15 of T-cells, Jensen's sarcoma, Kaposi's sarcoma, Kupffer cell sarcoma, angiosarcoma, leukosarcoma, malignant mesenchymoma sarcoma, parosteal sarcoma, reticulocytic sarcoma, Rous sarcoma, serocystic sarcoma, synovial sarcoma, and telangiectaltic sarcoma.

- The term "melanoma" is taken to mean a tumour arising from the melanocytic system of the skin and other organs. Melanomas include, for example, acral-lentiginous melanoma, amelanotic
- 20 melanoma, benign juvenile melanoma, Cloudman's melanoma, S91 melanoma, Harding-Passey melanoma, juvenile melanoma, lentigo maligna melanoma, malignant melanoma, nodular melanoma, subungal melanoma, and superficial spreading melanoma.

- The term "carcinoma" refers to a malignant new growth made up of epithelial cells tending to infiltrate the surrounding tissues and give rise to metastases. Exemplary carcinomas include, for
- 25 example, acinar carcinoma, acinous carcinoma, adenocystic carcinoma, adenoid cystic carcinoma, carcinoma adenomatosum, carcinoma of adrenal cortex, alveolar carcinoma, alveolar cell carcinoma, basal cell carcinoma, carcinoma basocellulare, basaloid carcinoma, basosquamous cell carcinoma, bronchioalveolar carcinoma, bronchiolar carcinoma, bronchogenic carcinoma, cerebriform carcinoma, cholangiocellular carcinoma, chorionic

carcinoma, colorectal carcinoma, colloid carcinoma, comedo carcinoma, corpus carcinoma, cribriform carcinoma, carcinoma en cuirasse, carcinoma cutaneum, cylindrical carcinoma, cylindrical cell carcinoma, duct carcinoma, carcinoma durum, embryonal carcinoma, encephaloid carcinoma, epierrmoid carcinoma, carcinoma epitheliale adenoides, exophytic carcinoma, carcinoma ex ulcere, carcinoma fibrosum, gelatiniform carcinoma, gelatinous carcinoma, giant cell carcinoma, carcinoma gigantocellulare, glandular carcinoma, granulosa cell carcinoma, hair-matrix carcinoma, haematoid carcinoma, hepatocellular carcinoma, Hurthle cell carcinoma, hyaline carcinoma, hypemephroid carcinoma, infantile embryonal carcinoma, carcinoma in situ, intraepidermal carcinoma, intraepithelial carcinoma, Krompecher's carcinoma, Kulchitzky-cell carcinoma, large-cell carcinoma, lenticular carcinoma, carcinoma lenticulare, lipomatous carcinoma, lymphoepithelial carcinoma, carcinoma medullare, medullary carcinoma, melanotic carcinoma, carcinoma molle, mucinous carcinoma, carcinoma muciparum, carcinoma mucocellulare, mucoepidermoid carcinoma, carcinoma mucosum, mucous carcinoma, carcinoma myxomatodes, naspharyngeal carcinoma, oat cell carcinoma, non-small cell carcinoma, carcinoma ossificans, osteoid carcinoma, papillary carcinoma, periportal carcinoma, preinvasive carcinoma, prickle cell carcinoma, pultaceous carcinoma, renal cell carcinoma of kidney, reserve cell carcinoma, carcinoma sarcomatodes, schneiderian carcinoma, scirrhus carcinoma, carcinoma scroti, signet-ring cell carcinoma, carcinoma simplex, small-cell carcinoma, solanoid carcinoma, spheroidal cell carcinoma, spindle cell carcinoma, carcinoma spongiosum, squamous carcinoma, squamous cell carcinoma, string carcinoma, carcinoma telangiectaticum, carcinoma telangiectodes, transitional cell carcinoma, carcinoma tuberosum, tuberous carcinoma, verrucous carcinoma, and carcinoma villosum.

The term "carcinoma" also encompasses adenocarcinomas. Adenocarcinomas are carcinomas that originate in cells that make organs which have glandular (secretory) properties or that originate in cells that line hollow viscera, such as the gastrointestinal tract or bronchial epithelia. Examples include, but are not limited to, adenocarcinomas of the breast, lung, colon, pancreas and prostate.

Additional cancers encompassed by the present invention include, for example, Hodgkin's Disease, Non-Hodgkin's lymphoma, multiple myeloma, neuroblastoma, rhabdomyosarcoma, primary thrombocytosis, primary macroglobulinemia, small-cell lung tumours, primary brain

tumours, malignant pancreatic insulanoma, malignant carcinoid, urinary bladder cancer, premalignant skin lesions, gliomas, testicular cancer, thyroid cancer, esophageal cancer, genitourinary tract cancer, malignant hypercalcemia, endometrial cancer, adrenal cortical cancer, mesothelioma and medulloblastoma.

- 5 The cancer to be treated or stabilized may be indolent or it may be aggressive. The compounds of the invention can be used to treat refractory cancers, advanced cancers, recurrent cancers and metastatic cancers. One skilled in the art will appreciate that many of these categories may overlap, for example, aggressive cancers are typically also metastatic.

- 10 “Aggressive cancer,” as used herein, refers to a rapidly growing cancer. One skilled in the art will appreciate that for some cancers, such as breast cancer or prostate cancer the term “aggressive cancer” will refer to an advanced cancer that has relapsed within approximately the earlier two-thirds of the spectrum of relapse times for a given cancer, whereas for other types of cancer, such as small cell lung carcinoma (SCLC), nearly all cases present rapidly growing cancers which are considered to be aggressive. The term can thus cover a subsection of a certain
- 15 cancer type or it may encompass all of other cancer types. A “refractory” cancer or tumour refers to a cancer or tumour that has not responded to treatment. “Advanced cancer,” refers to overt disease in a patient that is not amenable to cure by local modalities of treatment, such as surgery or radiotherapy. Advanced disease may refer to a locally advanced cancer or it may refer to metastatic cancer. The term “metastatic cancer” refers to cancer that has spread from one part of
- 20 the body to another.

The present invention also contemplates the use of the compounds of Formula I as “sensitizing agents,” which selectively inhibit the growth of cancer cells. In this case, the compound alone does not have a cytotoxic effect on the cancer cell, but provides a means of weakening the cancer cells, and thereby facilitates the benefit from conventional anti-cancer therapeutics.

- 25 Thus, the present invention contemplates the administration to a subject of a therapeutically effective amount of one or more compound of Formula I together with one or more anti-cancer therapeutics. The compound(s) can be administered before, during or after treatment with the anti-cancer therapeutic. An “anti-cancer therapeutic” is a compound, composition or treatment that prevents or delays the growth and/or metastasis of cancer cells. Such anti-cancer

therapeutics include, but are not limited to, chemotherapeutic drug treatment, radiation, gene therapy, hormonal manipulation, immunotherapy and antisense oligonucleotide therapy. Examples of useful chemotherapeutic drugs include, but are not limited to, hydroxyurea, busulphan, cisplatin, carboplatin, chlorambucil, melphalan, cyclophosphamide, Ifosphamide, danorubicin, doxorubicin, epirubicin, mitoxantrone, vincristine, vinblastine, Navelbine® (vinorelbine), etoposide, teniposide, paclitaxel, docetaxel, gemcitabine, cytosine, arabinoside, bleomycin, neocarzinostatin, suramin, taxol, mitomycin C and the like. The compounds of the invention are also suitable for use with standard combination therapies employing two or more chemotherapeutic agents. It is to be understood that anti-cancer therapeutics for use in the present invention also include novel compounds or treatments developed in the future.

VI. Pharmaceutical Compositions

The compounds of the present invention are typically formulated prior to administration. The present invention thus provides pharmaceutical compositions comprising one or more compounds of Formula I and a pharmaceutically acceptable carrier, diluent, or excipient. The pharmaceutical compositions are prepared by known procedures using well-known and readily available ingredients. Pharmaceutical compositions comprising one or more compounds of Formula I in combination with one or more known cancer chemotherapeutics are also contemplated by the present invention.

Compounds of the general Formula I or pharmaceutical compositions comprising the compounds may be administered orally, topically, parenterally, by inhalation or spray, or rectally in dosage unit formulations containing conventional non-toxic pharmaceutically acceptable carriers, adjuvants and vehicles. In the usual course of therapy, the active compound is incorporated into an acceptable vehicle to form a composition for topical administration to the affected area, such as hydrophobic or hydrophilic creams or lotions, or into a form suitable for oral, rectal or parenteral administration, such as syrups, elixirs, tablets, troches, lozenges, hard or soft capsules, pills, suppositories, oily or aqueous suspensions, dispersible powders or granules, emulsions, injectables, or solutions. The term parenteral as used herein includes subcutaneous injections, intradermal, intra-articular, intravenous, intramuscular, intravascular, intrasternal injection or infusion techniques.

Compositions intended for oral use may be prepared in either solid or fluid unit dosage forms. Fluid unit dosage form can be prepared according to procedures known in the art for the manufacture of pharmaceutical compositions and such compositions may contain one or more agents selected from the group consisting of sweetening agents, flavouring agents, colouring agents and preserving agents in order to provide pharmaceutically elegant and palatable preparations. An elixir is prepared by using a hydroalcoholic (e.g., ethanol) vehicle with suitable sweeteners such as sugar and saccharin, together with an aromatic flavoring agent. Suspensions can be prepared with an aqueous vehicle with the aid of a suspending agent such as acacia, tragacanth, methylcellulose and the like.

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- 10 Solid formulations such as tablets contain the active ingredient in admixture with non-toxic pharmaceutically acceptable excipients that are suitable for the manufacture of tablets. These excipients may be for example, inert diluents, such as calcium carbonate, sodium carbonate, lactose, calcium phosphate or sodium phosphate: granulating and disintegrating agents for example, corn starch, or alginic acid: binding agents, for example starch, gelatin or acacia, and
- 15 lubricating agents, for example magnesium stearate, stearic acid or talc and other conventional ingredients such as dicalcium phosphate, magnesium aluminum silicate, calcium sulfate, starch, lactose, methylcellulose, and functionally similar materials. The tablets may be uncoated or they may be coated by known techniques to delay disintegration and absorption in the gastrointestinal tract and thereby provide a sustained action over a longer period. For example, a time delay
- 20 material such as glyceryl monostearate or glyceryl distearate may be employed.

- Formulations for oral use may also be presented as hard gelatin capsules wherein the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules wherein the active ingredient is mixed with water or an oil medium, for example peanut oil, liquid paraffin or olive oil. Soft gelatin capsules are prepared by machine encapsulation of a slurry of the compound with an acceptable vegetable oil, light liquid petrolatum or other inert oil.
- 25

Aqueous suspensions contain active materials in admixture with excipients suitable for the manufacture of aqueous suspensions. Such excipients are suspending agents, for example sodium carboxymethylcellulose, methyl cellulose, hydropropylmethylcellulose, sodium alginate,

polyvinylpyrrolidone, gum tragacanth and gum acacia: dispersing or wetting agents may be a naturally-occurring phosphatide, for example, lecithin, or condensation products of an alkylene oxide with fatty acids, for example polyoxyethylene stearate, or condensation products of ethylene oxide with long chain aliphatic alcohols, for example hepta-decaethyleneoxycetanol, or
5 condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with partial esters derived from fatty acids and hexitol anhydrides, for example polyethylene sorbitan monooleate. The aqueous suspensions may also contain one or more preservatives, for example ethyl, or *n*-propyl- *p*-hydroxy benzoate, one or more colouring agents, one or more flavouring
10 agents or one or more sweetening agents, such as sucrose or saccharin.

Oily suspensions may be formulated by suspending the active ingredients in a vegetable oil, for example peanut oil, olive oil, sesame oil or coconut oil, or in a mineral oil such as liquid paraffin. The oily suspensions may contain a thickening agent, for example beeswax, hard paraffin or cetyl alcohol. Sweetening agents such as those set forth above, and flavouring agents may be
15 added to provide palatable oral preparations. These compositions may be preserved by the addition of an anti-oxidant such as ascorbic acid.

Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water provide the active ingredient in admixture with a dispersing or wetting agent, suspending agent and one or more preservatives. Suitable dispersing or wetting agents and
20 suspending agents are exemplified by those already mentioned above. Additional excipients, for example sweetening, flavouring and colouring agents, may also be present.

Pharmaceutical compositions of the invention may also be in the form of oil-in-water emulsions. The oil phase may be a vegetable oil, for example olive oil or peanut oil, or a mineral oil, for example liquid paraffin or mixtures of these. Suitable emulsifying agents may be naturally-
25 occurring gums, for example gum acacia or gum tragacanth, naturally-occurring phosphatides, for example soy bean, lecithin, and esters or partial esters derived from fatty acids and hexitol, anhydrides, for example sorbitan monooleate, and condensation products of the said partial esters with ethylene oxide, for example polyoxyethylene sorbitan monooleate. The emulsions may also contain sweetening and flavoring agents.

The pharmaceutical compositions may be in the form of a sterile injectable aqueous or oleaginous suspension. This suspension may be formulated according to known art using those suitable dispersing or wetting agents and suspending agents that have been mentioned above. The sterile injectable preparation may also be a sterile injectable solution or a suspension in a non-toxic parentally acceptable diluent or solvent, for example as a solution in 1,3-butanediol. Among the acceptable vehicles and solvents that may be employed are water, Ringer's solution and isotonic sodium chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose any bland fixed oil may be employed including synthetic mono- or diglycerides. In addition, fatty acids such as oleic acid find use in the preparation of injectables. Adjuvants such as local anaesthetics, preservatives and buffering agents can also be included in the injectable solution or suspension.

The compound(s) of the general Formula I may be administered, together or separately, in the form of suppositories for rectal administration of the drug. These compositions can be prepared by mixing the drug with a suitable non-irritating excipient which is solid at ordinary temperatures but liquid at the rectal temperature and will therefore melt in the rectum to release the drug. Such materials include cocoa butter and polyethylene glycols.

Other pharmaceutical compositions and methods of preparing pharmaceutical compositions are known in the art and are described, for example, in "*Remington: The Science and Practice of Pharmacy*" (formerly "*Remingtons Pharmaceutical Sciences*"); Gennaro, A., Lippincott, Williams & Wilkins, Philadelphia, PA (2000).

VII. Administration of Compounds of Formula I

Typically in the treatment of cancer, therapeutic compounds are administered systemically to a subject, for example, by bolus injection or continuous infusion into a subject's bloodstream. When used in conjunction with one or more known chemotherapeutic agents, the compounds can be administered prior to, or after, administration of the chemotherapeutic agents, or they can be administered concomitantly. The one or more chemotherapeutic may be administered systemically, for example, by bolus injection or continuous infusion, or it may be administered orally.

The compounds of Formula I may be used as part of a neo-adjuvant therapy (to primary therapy), as part of an adjuvant therapy regimen, where the intention is to cure the cancer in a subject. The present invention contemplates the use of the compounds of Formula I at various stages in tumour development and progression, including in the treatment of advanced and/or aggressive neoplasias (*i.e.* overt disease in a subject that is not amenable to cure by local modalities of treatment, such as surgery or radiotherapy), metastatic disease, locally advanced disease and/or refractory tumours (*i.e.* a cancer or tumour that has not responded to treatment).

“Primary therapy” refers to a first line of treatment upon the initial diagnosis of cancer in a subject. Exemplary primary therapies may involve surgery, a wide range of chemotherapies and radiotherapy. “Adjuvant therapy” refers to a therapy that follows a primary therapy and that is administered to subjects at risk of relapsing. Adjuvant systemic therapy is begun soon after primary therapy to delay recurrence, prolong survival or cure a subject.

It is contemplated that the compounds of the invention can be used alone or in combination with one or more other chemotherapeutic agents as part of an adjuvant therapy. Combinations of the compounds of Formula I and standard chemotherapeutics may act to improve the efficacy of the chemotherapeutic and, therefore, can be used to improve standard cancer therapies. This application is particularly important in the treatment of drug-resistant cancers which are not responsive to standard treatment. Drug-resistant cancers can arise, for example, from heterogeneity of tumour cell populations, alterations in response to chemotherapy and increased malignant potential. Such changes are often more pronounced at advanced stages of disease and have, in part, as an underlying cause, changes in genome/message stability.

The dosage to be administered is not subject to defined limits, but it will usually be an effective amount. It will usually be the equivalent, on a molar basis of the pharmacologically active free form produced from a dosage formulation upon the metabolic release of the active free drug to achieve its desired pharmacological and physiological effects. The compositions may be formulated in a unit dosage form. The term “unit dosage form” refers to physically discrete units suitable as unitary dosages for human subjects and other mammals, each unit containing a predetermined quantity of active material calculated to produce the desired therapeutic effect, in association with a suitable pharmaceutical excipient. Examples of ranges for the compound(s) in

each dosage unit are from about 0.05 to about 100 mg, or more usually, from about 1.0 to about 50 mg.

Daily dosages of the compounds of the present invention will typically fall within the range of about 0.01 to about 100 mg/kg of body weight, in single or divided dose. However, it will be understood that the actual amount of the compound(s) to be administered will be determined by a physician, in the light of the relevant circumstances, including the condition to be treated, the chosen route of administration, the actual compound administered, the age, weight, and response of the individual patient, and the severity of the patient's symptoms. The above dosage range is given by way of example only and is not intended to limit the scope of the invention in any way.

In some instances dosage levels below the lower limit of the aforesaid range may be more than adequate, while in other cases still larger doses may be employed without causing harmful side effects, for example, by first dividing the larger dose into several smaller doses for administration throughout the day.

VIII. Kits

The present invention additionally provides for therapeutic kits containing one or more compounds of Formula I for use in the treatment of cancer. The contents of the kit can be lyophilized and the kit can additionally contain a suitable solvent for reconstitution of the lyophilized components. Individual components of the kit would be packaged in separate containers and, associated with such containers, can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, for use or sale for human or animal administration.

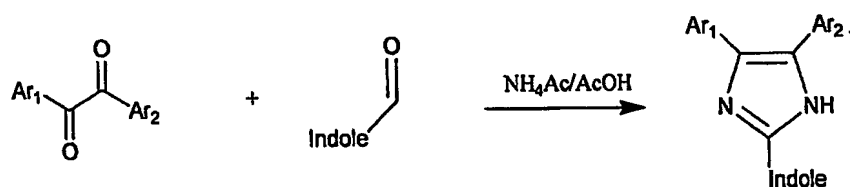
When the components of the kit are provided in one or more liquid solutions, the liquid solution can be an aqueous solution, for example a sterile aqueous solution. For *in vivo* use, the compounds may be formulated into a pharmaceutically acceptable syringeable composition. In this case the container means may itself be an inhalant, syringe, pipette, eye dropper, or other such like apparatus, from which the formulation may be applied to an infected area of the subject, such as the lungs, injected into an subject, or even applied to and mixed with the other components of the kit.

To gain a better understanding of the invention described herein, the following examples are set forth. It should be understood that these examples are for illustrative purposes only. Therefore, they should not limit the scope of this invention in any way.

EXAMPLES

5 Preparation of compounds:

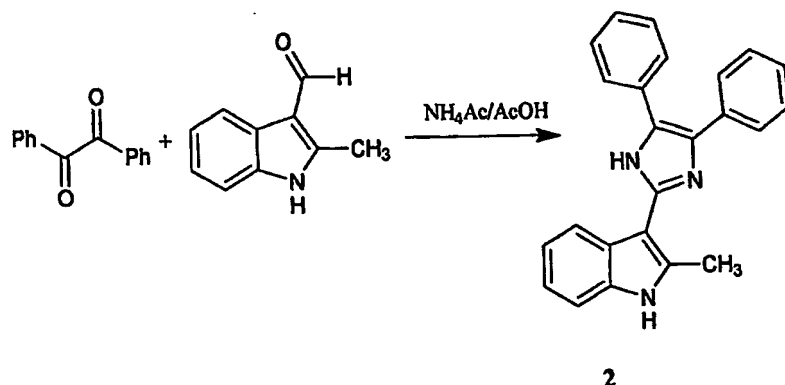
All reactions have been carried out according to the scheme shown below;



15 In a typical experimental procedure 1 mmol (1 equiv.) of the indole carboxyaldehyde was combined with 1.05 - 1.10 mmole (1.05 - 1.1 equiv.) of the benzil and 20 mmole (20 equiv.) of ammonium acetate and 5 ml of acetic acid. The mixture was magnetically stirred and heated to reflux for 3-5 hr. The reaction process was monitored by TLC, until complete consumption of the indole was achieved. The reaction mixture was cooled to room temperature and added drop-wise into well-stirred ice-water. The suspension solid was then filtered and the crude solid was dissolved in ethyl acetate, dried over sodium sulfate and filtered, the organic solvent was removed by vacuum. The products was then either recrystallized with alcohol or separated by column chromatography using petroleum ether-Ethyl acetate as an eluant.

25 It is noteworthy that, the TLC of the products shows a characteristic blue florescent color under the UV (wave length $\lambda = 254\text{nm}$), a property used as an additional characterization feature. Melting points were recorded using a MEL-TEMP capillary melting point apparatus, the melting point are uncorrected. ¹H-NMR was performed in a 500 MHz Bruker instrument at room temperature using a suitable deuterated solvent.

Example 1: Preparation of compound 2



¹H-NMR: δ (DMSO- d_6), 12.10 (s, 1H), 11.30 (s, 1H), 7.98 (d, 1H), 7.62 (d, 2H), 7.56 (d, 2H),
 5 7.45 (t, 2H), 7.28-7.40 (m, 4H), 7.24 (t, 1H), 7.03-7.14 (m, 2H), 2.70 (s, 3H). HRMS m/z for $C_{24}H_{19}N_3$ calc. Is 349.157898, found 349.157897. M.p.= decomposed at 260-264.

Example 2: Compound 5

¹H-NMR ($CDCl_3$): δ = 8.02 (d, 2H), 7.53 (d, 1H), 7.43 - 7.52 (m, 6H), 7.41 (d, 1H), 7.21 - 7.34
 10 (m, 6H), 2.81 (s, 3H), 2.75 (s, 3H). EIMS [M^+] m/z for $C_{28}H_{23}N_3O_2$ is 433. M.p.= 224-227.

Example 3: Compound 11

¹H-NMR ($CDCl_3$): δ = 7.70 (d, 1H), 7.41 (d, 4H), 7.32 (d, 1H), 7.09 (q, 2H), 6.77 (d, 4H), 2.95
 15 (s, 12H), 2.67 (s, 3H). EIMS [M^+] m/z for $C_{28}H_{29}N_5$ is 435, M.p.= decomposed at 236-238.

Example 4: Compound 13

¹H-NMR ($CDCl_3$): δ = 7.47 (d, 4H), 7.44 (d, 4H), 7.30-7.34 (m, 1H), 7.14 - 7.19 (m, 3H), 2.68
 (bs, 3H), EIMS [M^+] m/z for $C_{24}H_{17}N_3Br_2$ is 507. M.p.= 240-245.

Example 5: Compound 19

¹H-NMR (DMSO- d_6): δ = 12.13 (s, 1H), 11.33 (s, 1H), 7.94 (d, 2H), 7.57 (d, 2H), 7.52 (bd, 2H),
 7.39 (bd, 2H), 7.35 (d, 1H), 7.05-7.12 (m, 3H), 2.50 (s, 3H). EIMS [M^+] m/z for $C_{24}H_{17}N_3Cl_2$ is
 418. M.p.= 165-167.

Example 6: Compound 29

¹H-NMR (DMSO-d₆): δ = 8.83 (q, 2H), 8.73 (m, 1H), 8.68 (d, 1H), 8.46 (d, 1H), 8.24 (s, 1H), 7.74 (t, 2H), 7.62 (t, 2H), 7.51 - 7.56 (m, 1H), 7.23 - 7.27 (m, 2H), 2.71 (s, 3H). EIMS [M⁺] m/z for C₂₃H₁₅N₃ is 303. M.p.= 135-137.

5

Example 7: Compound 31

¹H-NMR (CDCl₃): δ = 8.90 (bs, 1H), 7.62 (bs, 1H), 7.48 (bd, 4H), 7.34 (m, 4H), 7.21 (m, 1H), 7.13 (m, 2H), 2.43 (bs, 3H). EIMS [M⁺] m/z for C₂₃H₁₅N₃ClBr is 448. M.p.= decomposed at 218-220.

10

Example 8: Compound 35

¹H-NMR (CDCl₃): δ = 7.78 (bs, 1H), 7.59 (d, 2H), 7.54 (d, 2H), 7.35 - 7.39 (m, 2H), 7.28 - 7.34 (m, 2H), 7.13 - 7.18 (m, 2H), 7.01 - 7.05 (m, 2H), 2.72 (bs, 3H). EIMS [M⁺] m/z for C₂₄H₁₈N₃F is 367. M.p.= decomposed at 247-250.

15

Example 9: Compound 38

¹H-NMR (Acetone-d₆): δ = 11.12 (bs, 1H), 10.46 (bs, 1H), 8.12 (d, 1H), 7.80 (bd, 2H), 7.62 (bd, 2H), 7.38-7.48 (m, 5H), 6.98 - 7.22 (m, 12H), 2.84 (bs, 3H). EIMS [M⁺] m/z for C₃₆H₂₇N₃O₂ is 533. M.p.= decomposed at 128-130.

20

Example 10: Compound 40

¹H-NMR (CDCl₃): δ = 8.12 (dd, 2H), 7.60 (m, 6H), 7.24-7.53 (m, 10H), 6.87 (bd, 2H), 6.61 (bd, 2H), 2.08 (s, 3H). M.p.= decomposed at 142.

Example 11: Compound 42

¹H-NMR (CDCl₃): δ = 9.39 (bs, 1H), 7.39 - 7.50 (m, 4H), 7.28 - 7.38 (m, 6H), 7.06 (bs, 1H), 6.94 (bs, 2H), 2.08 (bs, 3H). EIMS [M⁺] m/z for C₂₄H₁₈N₃Br is 428. M.p.= decomposed at 155-157.

Example 12: Compound 44

$^1\text{H-NMR}$ (CDCl_3): δ = 8.78 (dd, 2H), 8.19 (dd, 1H), 7.96 (bs, 1H), 7.80 (dd, 1H), 7.80 (dd, 1H), 7.55-7.77 (m, 6H), 7.16-7.42 (m, 2H), 2.87 (bs, 3H). EIMS [M^+] m/z for $\text{C}_{24}\text{H}_{17}\text{N}_3$ is 347. M.p.= decomposed at 167.

5 **Example 13: Compound 10**

$^1\text{H-NMR}$ (CDCl_3): δ = 10.68 (bs, 1H), 7.73 (bs, 1H), 7.22 (d, 4H), 6.99 (bs, 1H), 6.92 (bd, 2H), 6.85 (bd, 2H), 6.611 (d, 4H), 3.70 (s, 6H). EIMS [M^+] m/z for $\text{C}_{25}\text{H}_{20}\text{N}_3\text{BrO}_2$ is 474. M.p.= 135.

10 **Example 14: Compound 26**

$^1\text{H-NMR}$ (DMSO-d_6): δ = 12.60 (s, 1H), 11.70 (s, 1H), 8.60 (d, 1H), 8.17 (s, 1H), 7.68 (bs, 1H), 7.46 (d, 2H), 7.33 (d, 2H), 7.25 (bs, 2H), 7.09 (bs, 1H). EIMS [M^+] m/z for $\text{C}_{19}\text{H}_{12}\text{N}_3\text{BrS}_2$ is 426.

15

Example 15: Compound 28

$^1\text{H-NMR}$ (DMSO-d_6): δ = 12.60 (s, 1H), 11.65 (s, 1H), 8.44- 8.64 (m, 3H), 8.01- 8.14 (m, 1H), 7.22-7.66 (m, 8H). EIMS [M^+] m/z for $\text{C}_{22}\text{H}_{14}\text{N}_4\text{BrF}$ is 433. M.p.= decomposed at 343.

20 **Example 16: Compound 32**

$^1\text{H-NMR}$ (CDCl_3): δ = 8.12 (bs, 1H), 7.48 (d, 2H), 7.46 (d, 2H), 7.23-7.34 (m, 8H). M.p.=230-232.

Example 17: Compound 34

25 $^1\text{H-NMR}$ (DMSO-d_6): δ = 12.63 (s, 1H), 11.67 (s, 1H), 8.62 (d, 1H), 8.21 (d, 2H), 8.08 (d, 1H), 7.86 (d, 2H), 7.39-7.64 (m, 6H), 7.32 (dd, 1H). EIMS [M^+] m/z for $\text{C}_{23}\text{H}_{15}\text{N}_4\text{BrFO}_2$ is 459. M.p.= decomposed at 250-253.

Example 18: Compound 36

30 $^1\text{H-NMR}$ (CDCl_3): δ = 10.42 (bs, 1H), 7.86 (s, 1H), 7.16-7.33 (m, 6H), 7.04 (dd, 2H), 6.95 (dd, 2H), 6.88 (t, 3H). EIMS [M^+] m/z for $\text{C}_{23}\text{H}_{15}\text{N}_3\text{BrF}$ is 432. M.p.= decomposed at 83-86.

Example 19: Compound 41

¹H-NMR (CDCl₃): δ = 8.08 (d, 4H), 8.07 (bs, 1H), 7.75 (d, 4H), 7.28-7.50 (m, 10H), 7.12 (bd, 2H), 6.97 (bs, 1H). M.p.= 155-158.

5

Example 20: Compound 43

¹H-NMR (CDCl₃): δ = 9.75 (bs, 1H), 7.83 (bs, 1H), 7.36 (m, 3H), 7.25 - 7.29 (m, 5H), 7.12 (m, 3H), 7.10 (bd, 1H). EIMS [M⁺] m/z for C₂₃H₁₅N₃Br₂ is 493. M.p.= decomposed at 230.

10 **Example 21: Compound 45**

¹H-NMR (DMSO-d₆): δ = 13.30 (bs, 1H), 11.62 (d, 1H), 8.87 (bd, 2H), 8.64 (bs, 1H), 8.44 (bs, 1H), 8.29 (t, 1H), 7.76 (t, 2H), 7.62 (t, 2H), 7.52 (d, 1H), 7.35 - 7.41 (m, 2H). EIMS [M⁺] m/z for C₂₃H₁₄N₃Br is 412.

15 **Example 22: Compound 84**

¹H-NMR (DMSO-d₆): δ = 8.64 (d, 1H), 8.17 (d, 1H), 7.47 (d, 1H), 7.39 (t, 1H), 7.33 (dd, 1H), 7.20-7.31 (m, 2H), 7.12 (bd, 2H), 6.97 (bd, 1H), 6.84 (bd, 1H). 3.77 (s, 3H), 3.72 (s, 3H), EIMS [M⁺] m/z for C₂₅H₂₀N₃BrO₂ is 474. M.p.= decomposed at 250-253.

20 **Example 23: Compound 37**

¹H-NMR (CDCl₃): δ = 9.92 (bs, 1H), 8.17 (bs, 1H), 7.87 (t, 1H), 7.55 (bs, 1H), 7.21-7.33 (m, 6H), 7.15-7.2 (m, 1H), 7.04-7.07 (m, 2H), 6.90 (t, 2H)). EIMS [M⁺] m/z for C₂₃H₁₆N₃F is 353. M.p.= 51.

25 **Example 24: Compound 46**

¹H-NMR (DMSO-d₆): δ = 13.09 (s, 1H), 11.61 (d, 1H), 8.83 (q, 2H), 8.73 (m, 1H), 8.68 (d, 1H), 8.46 (d, 1H), 8.24 (s, 1H), 7.74 (t, 2H), 7.62 (t, 2H), 7.51 - 7.56 (m, 1H), 7.23 - 7.27 (m, 2H). EIMS [M⁺] m/z for C₂₃H₁₅N₃ is 333. M.p.= 135-137.

30 **Example 25: Compound 83**

¹H-NMR (DMSO-d₆): δ = 12.40 (s, 1H), 11.80 (s, 1H), 8.56 (d, 1H), 8.24 (s, 1H), 8.15 (s, 1H), 7.78 (d, 1H), 7.65 (d, 2H), 7.54 (d, 2H), 7.47 (t, 2H), 7.32 - 7.42 (m, 3H), 7.24 (t, 1H), 3.90 (s, 3H). EIMS [M⁺] m/z for C₂₅H₁₉N₃O₂ is 393. M.p.= 293-295.

5 **Example 26: Compound 74**

¹H-NMR (DMSO-d₆): δ = 11.74 (d, 1H), 7.95 (dd, 1H), 7.32-7.57 (m, 6H), 7.17-7.31 (m, 2H), 7.12 (t, 1H), 6.70 (d, 1H), 6.66 (bd, 1H). EIMS [M⁺] m/z for C₂₃H₁₉N₂FO₂ is 374.

Example 27: *In vitro* Inhibition of Proliferation of Cancer Cells

10 Selected compounds of Formula I were tested for anti-cancer activity *in vitro* using a human colon carcinoma cells (HT-29) and human non-small cell lung cancer cells (H460). The cells were maintained in α-MEM medium (Wisent, St-Bruno, Qc) supplemented with 10% FBS, and grown at 37°C in an atmosphere of 5% CO₂. Cells were transferred onto 150mm tissue culture plates and grown until sub-confluency (70-80%) prior to their use.

15 The anti-cancer activity *in vitro* was evaluated by a cell proliferation assay based on the ability of live cells to reduce the tetrazolium salt XTT to orange colored compounds of formazan (XTT cell proliferation kit II, Roche Applied Science, Montreal, QC).

20 Approximately 4 x 10³ colon cancer cells (HT-29) or 2 x 10³ non-small cell lung cancer cells (NCI-H460) in 100 µl of complete culture medium were plated onto 96-well microtiter plates and incubated overnight at 37°C. The medium was then removed by inverting the plate and patting on a sterile absorbent cloth. Fifty µl of medium containing the test compound at either 25 or 100 µM, were added to the wells containing cells and incubated at 37°C in an atmosphere of 5% CO₂ for 48 h. Following incubation, 25 µl of an XTT reaction mixture (XTT at a final concentration of 0.3 mg/ml) were added to each well and the plates were incubated for a further 4 h. The absorbance of each sample was then determined at a wavelength of 490 nm/650 nm as
25 reference. Each compound was tested in duplicate and the results are reported as averages. Table II shows the effect that different compounds of Formula I have on the growth of human colon carcinoma HT-29. Table III shows the effect that different compounds of Formula I have on the growth of human non-small cell lung cancer cells (H460).

Table II: Inhibition of Proliferation of Human Colon Carcinoma (HT-29) Cells

| Compound | 100 μ M | | 25 μ M | |
|----------|-------------|-------|------------|-------|
| | % Survival | SD(%) | % Survival | SD(%) |
| 5 | 110.7 | 1.9 | 110.9 | 2.8 |
| 6 | 3.2 | 0.2 | 11.7 | 1.6 |
| 9 | 15.1 | 2.8 | 68.3 | 16 |
| 10 | 7.6 | 0.5 | 25.8 | 2.6 |
| 11 | 94.3 | 3.6 | 107.8 | 1 |
| 13 | 82.4 | 0.8 | 105.9 | 5.4 |
| 14 | 3.8 | 0.5 | 55.2 | 15.7 |
| 19 | 37.1 | 7.2 | 105.5 | 2.9 |
| 20 | 28.1 | 5 | 100 | 2.7 |
| 23 | 45.7 | 5.8 | 98.2 | 0 |
| 25 | 39.8 | 4.7 | 63.9 | 1.6 |
| 27 | 35 | 0.6 | 62.3 | 2.4 |
| 29 | 20.9 | 1 | 37.1 | 6.2 |
| 31 | 24.9 | 1.8 | 98.6 | 3.3 |
| 32 | 7.7 | 0.6 | 22.7 | 0 |
| 33 | 10 | 0.3 | 56.1 | 5.9 |
| 34 | 10.8 | 0 | 22.8 | 2 |
| 35 | 2.5 | 0.3 | 44.1 | 4.4 |
| 36 | 4.7 | 0.8 | 31.6 | 2.2 |
| 38 | 35.7 | 1.5 | 67 | 7.9 |
| 39 | 53 | 0.3 | 96.2 | 4 |
| 40 | 36.7 | 1.8 | 79.1 | 1.4 |
| 42 | 1.8 | 0 | 59.9 | 0.1 |
| 43 | 5.7 | 0.3 | 28 | 6.8 |
| 44 | 6.5 | 0.4 | 63.6 | 2.9 |
| 45 | 35 | 0.6 | 88.9 | 3.3 |
| 46 | 4.5 | 0 | 16.1 | 1 |

| Compound | 100µM | | 25µM | |
|----------|------------|-------|------------|-------|
| | % Survival | SD(%) | % Survival | SD(%) |
| 73 | 62.2 | 3.2 | 65 | 2 |
| 83 | 109.5 | 4.7 | 100.3 | 1.1 |
| CPT-11 | 51.1 | 3.2 | 82.3 | 10 |
| Vehicle | 100 | 7 | 100 | 7 |

Table III: Inhibition of Proliferation of Human Lung Carcinoma (NCI- 460) Cells

| Compound | 100µM | | 25µM | |
|----------|------------|-------|------------|-------|
| | % Survival | SD(%) | % Survival | SD(%) |
| 5 | 106 | 2.6 | 102 | 0.7 |
| 6 | 1.9 | 0.5 | 10.4 | 0.8 |
| 9 | 8.4 | 1.6 | 98.2 | 1 |
| 10 | 2.7 | 0.1 | 26.9 | 1.6 |
| 11 | 101.6 | 8.3 | 98.8 | 3.3 |
| 13 | 96.2 | 1.1 | 101.9 | 4 |
| 14 | 1.8 | 0.1 | 83.5 | 20.4 |
| 19 | 27.3 | 6.1 | 89.2 | 0.2 |
| 20 | 82.1 | 20.6 | 98.6 | 2.1 |
| 23 | 92.1 | 0 | 96.3 | 0.9 |
| 25 | 89 | 3.4 | 99.5 | 0.8 |
| 27 | 43.1 | 1.4 | 93.5 | 0.2 |
| 29 | 20.2 | 1.4 | 73.8 | 2.2 |
| 31 | 37.6 | 5.6 | 94 | 2.3 |
| 32 | 2.9 | 0.5 | 15 | 0.2 |
| 33 | 9.4 | 2 | 73.4 | 2.2 |
| 34 | 7.6 | 0.9 | 17.4 | 0.2 |
| 35 | 1.2 | 0.1 | 83.8 | 8 |
| 36 | 2.4 | 0.3 | 24.5 | 2.1 |

| Compound | 100µM | | 25µM | |
|----------|------------|-------|------------|-------|
| | % Survival | SD(%) | % Survival | SD(%) |
| 38 | 12.2 | 1.6 | 98.8 | 2 |
| 39 | 17 | 0.7 | 98.1 | 0.3 |
| 40 | 7.7 | 0.5 | 97.6 | 3 |
| 42 | 1.2 | 0.1 | 66.1 | 16.9 |
| 43 | 3 | 0.1 | 18.8 | 1.5 |
| 44 | 3.4 | 1 | 77.3 | 5.3 |
| 45 | 32.7 | 5.5 | 96 | 1.1 |
| 46 | 1.9 | 0.1 | 11.5 | 1.4 |
| 73 | 53.5 | 1.3 | 89.2 | 0.6 |
| 83 | 109.2 | 1.5 | 100.9 | 2.6 |
| CPT-11 | 6.1 | 0.5 | 32.2 | 4.5 |
| Vehicle | 100 | 3.4 | 100 | 3.4 |

Example 28: *In Vivo* Inhibition of Cancer Cell Growth

In vivo efficacy studies of various compounds of Formula I were also performed using a mouse xenograft model using the human colon adenocarcinoma cell line HT-29. CD-1 female nude mice (6-7 weeks) were divided into groups of five and each injected subcutaneously in the lower mid-back with HT-29 cells (3×10^6 cells in 0.1 ml PBS). Treatment was initiated 5 days post-inoculation (size of tumours = 20-40 mm³). The treatment schedule consisted of 2 x 200 µl intraperitoneal injections per day of 5 mg/ml (100 mg/Kg/d) of test compound for five days a week, for 4 weeks. Tumour sizes were measured during the course of the treatment using calipers. After 4 weeks, the mice were sacrificed by cervical dislocation and the tumours were surgically removed and weighed. Figure 1 shows the average tumour size (mm³) in the different groups of mice. Figures 2 and 3 show the average tumour weight per group of mice and per individual mouse, respectively.

Example 29: Selectivity of Compounds of Formula I

Compounds of Formula I that demonstrate the ability to decrease the growth or proliferation of at least one cancer cell line may undergo further testing to evaluate their selectivity towards cancer cells. An exemplary method to measure the selectivity of the compounds of the present invention is provided below.

IC₉₀ values of selected compounds on a panel of normal actively proliferating cells (HUVEC and WI38) and cancer cells representing colon (HT-29), lung (NCI-H460), breast (MDA-MB-231) and prostate cancer (PC-3) are measured. Compounds with 2-fold or higher overall selectivity to the panel of cancer cell lines at IC₉₀ are identified as potential therapeutics.

IC₉₀ values are determined using the XTT assay as an indicator of growth arrest and/or cytotoxicity. This assay is conducted as outlined in Example 27. Percentage inhibition is calculated for each cell line and IC₉₀ values for each compound and cell type determined. The average IC₉₀ values for the normal cells are calculated and divided by the average IC₉₀ values for the cancer cell lines. Compounds with a selectivity ratio of >2 are identified and chosen for further optimization and/or testing.

Example 30: *In vitro* IC₅₀ and GI₅₀ Evaluations against Panels of Cancer Cell Lines

Compounds of Formula I that demonstrate the ability to decrease the growth or proliferation of at least one cancer cell line may undergo further testing to determine whether they possess significant potency across a broader range of cancer cell types and to identify whether the potency reveals cell-type or genotype specificity. Such studies may provide important clues as to potential mechanisms of action by these compounds.

Selected compounds are evaluated against a panel of cancer cell lines, for example cell lines representing common cancers such as colon (e.g. HT-29, COLO320, HCT-116), breast (e.g. MDA-MB-231, T47D), prostate (e.g. PC-3, DU145), lung (e.g. NCI-H460, H209), bladder (e.g. T24), brain (e.g. U-87-MG), ovary (e.g. SK-OV-3), pancreas (e.g. AsPC-1, SU.86.86, BxPC-3), kidney (e.g. A498, Caki-1), liver (e.g. HepG2) and skin (e.g. A2058, C8161) cancer. IC₅₀ values are determined using XTT assays as described above. Correlative studies are carried out to investigate whether the potency estimated by IC₅₀ determinations demonstrates a clear

dependency on tumour cell types or the status of mutations in such genes involved in cell-cycle check point controls (e.g. p53 and pRb).

In addition, if desired, the selected compounds can be submitted to a independent testing facility, such as the U.S. National Cancer Institute (NCI) of the National Institutes of Health (NIH) in Rockwell, Maryland, for evaluation of their anti-proliferative activity. The NCI conducts a standard 48/72 hour 60 cell line assay and an *in vitro* time course assay as described in Alley *et al.* (*Cancer Res* (1988) 48:589). In the standard 60 cell line assay, a minimum of 5 concentrations of the test compound are tested at 10-fold dilutions against 60 cell lines and cell growth is assayed at 48 and 72 hours using a sulphorhodamine B assay. For the time course analysis, tumour cells are treated with the test compound at various time points, then washed and grown in medium free of the test compound until the end of the experiment at 144 hrs. This assay employs 20% FBS to better approximate the minimum c x t (concentrations and times) test compound exposure conditions that are required to achieve activity *in vivo*. Cell growth is quantified by an MTT assay (similar to the XTT assay described above) and the concentration of the test compound required for growth inhibition is determined. The inhibitory effect of the test compounds are expressed as a GI₅₀ value, which represents the molar concentration of the test compound that results in 50% growth inhibition.

Example 31: *In Vivo* Anti-tumour Efficacy Evaluations

Pharmacological evaluation of selected compounds is conducted in animal models of human tumour growth. Data from these studies provide evidence of the therapeutic efficacy of selected compounds against various types of cancer and help to identify compounds with better pharmacological properties and potency.

Examples of mouse models that can be utilized to investigate the efficacy of selected compounds include, but are not limited to, xenografts of various human tumour types, inoculated subcutaneously into nude mice or mice with severe combined immunodeficiency disorder (SCID); orthotopic implantation of various human tumours in nude or SCID mice for investigation of effects on the tumour in the target organ (for example, a pancreatic cancer cell graft implanted directly into the pancreas of the animal), and investigation of spontaneous tumours in normal mice.

In order to provide evidence of the efficacy of a selected compound as a single agent, it may be evaluated, for example, in specific models (xenograft or orthotopic) for representative human cancers such as pancreas, skin (melanoma), kidney, colon, breast, lung, liver, ovary, prostate, bladder and brain. Similar studies can be conducted to evaluate the performance of test compounds in combination with other standard therapeutic modalities used in the treatment of human cancers.

For typical xenograft studies, 5-6 week old, female, CD-1 athymic nude mice, (Charles River, Montreal, QC) are acclimatized in a pathogen-free facility for at least 1 week. Animal protocols followed are in compliance with the Guide for the Care and Use of Laboratory Animals in Canada. Approximately 10^6 - 10^7 human tumour cells in 100 ml PBS are subcutaneously injected into the right flank of each mouse. Once tumours reach an approximate volume of 100 mm^3 (several days post tumour cell injection), mice are randomized by tumour size into control and treatment groups. Test compounds are administered at various doses 5 days a week for several weeks. Control animals receive vehicle alone (negative control) and/or a standard chemotherapeutic (positive control) for the same period. The tumour dimensions (length, width, and height) are measured using calipers twice a week over the treatment period. Tumour volume is calculated by the formula $L \times W \times H/2$, where L indicates length, W indicates width and H indicates height. The mice are sacrificed when the tumour burden reaches approximately 10% of total body weight and excised tumours are weighed. A standard bar graph is used to demonstrate the differences in tumour weights with each bar representing mean tumour weight.

Example 32: Assays to Investigate Potential Mechanism of Action

The potential mechanism of action of selected compounds can be investigated using assays such as cell-cycle analysis, apoptosis assays, anti-angiogenesis assays and immunohistochemical analysis. A representative example of each type of assay is provided below.

25 i) *Cell-cycle analysis*

Alterations in cell cycle are determined using flow cytometric analyses. Tumour cells sensitive to a test compound are synchronized by plating in medium containing 0.5% FBS for 24 h followed by culturing in FBS-free medium for 48 h. The cells are then released into complete

medium containing 0.1% DMSO (vehicle control) or the test compound at an appropriate concentration (e.g. $3 \times \text{IC}_{90}$ value), harvested 16 to 24 h following treatment, washed twice with cold PBS and fixed in 70% ethanol at 4° C for at least 4 h. The fixed cells are centrifuged at 1500rpm for 4 minute at 4° C, washed twice with cold PBS containing 2% FBS, treated with 3 mg/ml ribonuclease (Sigma Chemical Co. Oakville, ON) and 50 µg/ml propidium iodide (PI) (Sigma Chemical Co.) for 30 minutes at 37° C. The fluorescence of the stained cells is measured using a FACScan flow cytometer and the Cell Quest program (Becton Dickinson, San Jose, CA). Data are evaluated using Modfit software (Verity software House, Topsham, ME) and the effects of the selected compounds on cell cycle are evaluated.

10 ii) *Apoptosis assay*

DNA fragmentation analysis is used to evaluate the apoptotic effects of test compounds. Briefly, cells are plated in six-well culture plates 24 hr prior to treatment. After incubation with the test compound, medium containing detached cells is transferred to 15 ml conical tubes while cells still attached to the plate are trypsinized and then added to the same tubes. After centrifugation, collected cells are washed with PBS and resuspended in 0.5 ml lysis buffer containing 50 mM Tris-HCl, pH 8.0, 1.0 M NaCl, 10 mM EDTA and 0.5% SDS. Cell lysates are transferred to microfuge tubes and proteinase K is added to a final concentration of 0.2 ml/ml and incubated overnight at 37° C. DNA is extracted by phenol:chloroform:isoamyl alcohol (24:24:1), dried and dissolved in 40 µM of 10 µM Tris-HCl (pH 8.0) and 0.1 mM EDTA. DNase-free RNase A is added to each sample for 30 min at 37° C and 12 µl of each sample are loaded onto a 2% agarose gel containing 0.5 µg/ml ethidium bromide and electrophoresed. DNA is visualized under UV illumination and the induction of apoptosis by the test compound is evaluated based on the generation of a nucleosomal-size DNA ladder.

iii) *Anti-angiogenesis assay*

25 Proliferation of new capillaries, i.e. angiogenesis or neovascularization, is critical for the transition of a small localized tumour to expand into a large malignant growth. The Matrigel Plug Assay (see, Passaniti *et al.*, *Lab. Invest.* (1992) 67:519) is a simple method for assessing angiogenesis and the possible anti-angiogenic effect of selected compounds in mice. Briefly, liquid Matrigel (Becton Dickinson & Co., NJ) is injected subcutaneously near the abdominal

midline or the dorsal flank of the animal using a 25-gauge needle. Growth factor-reduced Matrigel supplemented with 8.3 nM basic fibroblast growth factor (bFGF, Collaborative Biomedical Products, MA) stays in liquid form at 4 °C. bFGF is a proven and potent inducer of angiogenesis. When injected into a mouse (0.5 ml/mouse), Matrigel immediately forms a readily recoverable solid gel, which is removed at various times (not exceeding 10 days) to assess neo-vessel growth around and into the gel. Test compounds are administered according to appropriate doses and schedules. Typically at a 5-day point, mice are sacrificed, overlying skin is removed and the gels are cut out retaining the peritoneal lining for support. For quantitation of angiogenesis, two methods are employed: 1. haemoglobin content in the gel is measured using the Drabkin method (Drabkin and Austin, *J. Biol Chem.* (1932) 98:719) and Drabkin reagent kit 525 (Sigma, MO); 2. the number of blood vessels invading the Matrigel is determined by microscopic analysis after the gels are fixed, embedded in paraffin, sectioned and stained.

iv) *Immunohistochemistry*

The anti-cancer effects of test compounds can be evaluated in mouse xenograft models (as described above) by quantitating the effects of these compounds on tumour growth, differentiation, apoptosis and angiogenesis using immunohistochemical methods.

Tumour cell proliferation, angiogenesis and tumour immune infiltrates are delineated immunohistochemically using specific antibodies (Ki-67 for proliferation, CD31 for angiogenesis and NK1.1 for NK cells and F4/80 for macrophage). Apoptosis is delineated utilising the TUNEL assay (*In Situ* Cell Death Detection kit; Boehringer Mannheim, Laval, QC). Signal generation is accomplished by peroxidase catalyzed generation of enzyme product which is visualized microscopically. Tissue histology is determined after H&E staining of separate sections.

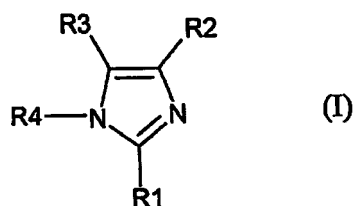
Briefly, tumour xenografts from treated mice are isolated, fixed and paraffin embedded individually in blocks and several 5 µm sections are cut for immunostaining and TUNEL assays. One additional section is obtained for H&E staining. For all immunohistochemical labeling, prior antigen retrieval is employed to improve detection. Typically, a 3-step amplification method is used to generate signals in immunohistochemistry that consists essentially of applying

a biotinylated secondary antibody that recognizes the primary antibody, followed by avidin-
peroxidase incubation. The final step is enzyme reaction in stable DAB solution.
Immunohistochemical sections are counterstained with hematoxylin for tissue histology. To
eliminate non-specific immunostaining with mouse monoclonal antibodies applied to mouse
5 tissues, a specific blocking step is included in the procedure. Staining patterns are documented
photographically, examined by at least two independent observers and quantitated by counting a
pre-determined number of cells.

The invention being thus described, it will be obvious that the same may be varied in many
ways. Such variations are not to be regarded as a departure from the spirit and scope of the
10 invention, and all such modifications as would be obvious to one skilled in the art are intended to
be included within the scope of the following claims.

THE EMBODIMENTS OF THE INVENTION IN WHICH AN EXCLUSIVE PROPERTY OR PRIVILEGE IS CLAIMED ARE DEFINED AS FOLLOWS:

1. A method of inhibiting neoplastic cell growth in a mammal comprising administering to said mammal a therapeutically effective amount of a compound of formula I,



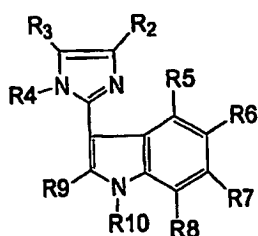
wherein:

R1 is aryl, substituted aryl, heterocycle, substituted heterocycle, heteroaryl, or substituted heteroaryl;

R2 and R3 are independently aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, or substituted heteroaryl or R2 and R3 when taken together along with the carbon atoms they are attached to, form aryl or substituted aryl, and

R4 is hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano.

2. The method according to claim 1, wherein said compound of structural formula I is administered in combination with one or more anti-cancer agent(s).
3. A method of inhibiting neoplastic cell growth in a mammal comprising administering to said mammal a therapeutically effective amount of a compound of formula II:



II

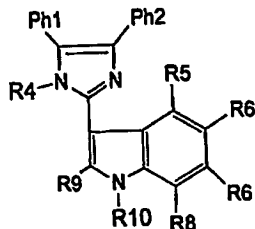
or a salt thereof, wherein:

R2 and R3 are independently aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, or substituted heteroaryl or R2 and R3 when taken together along with the carbon atoms they are attached form aryl or substituted aryl, heterocycle, heteroaryl, substituted heterocycle, or substituted heteroaryl;

R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano;

R10 is H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, acyl, -CH₂-aryl, -CH₂-heteroaryl.

4. A method of inhibiting neoplastic cell growth in a mammal comprising administering to said mammal a therapeutically effective amount of a compound of formula III:



III

or a salt thereof, wherein:

Ph1 and Ph2 are independently selected from phenyl and substituted phenyl;

R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano;

R10 is H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, acyl;

with the proviso that the compounds are other than:

3-(4,5-diphenyl-1H-imidazol-2-yl)-1-methyl-1H-indole;

3-[4-(4-chlorophenyl)-5-phenyl-1H-imidazol-2-yl]-1-methyl-1H-indole;

3-[4-(4-bromophenyl)-5-phenyl-1H-imidazol-2-yl]-1-methyl-1H-indole;

3-[4-(4-methylphenyl)-5-phenyl-1H-imidazol-2-yl]-1-methyl-1H-indole;

3-[4-(4-methoxyphenyl)-5-phenyl-1H-imidazol-2-yl]-1-methyl-1H-indole;

3-[4-(4-ethoxyphenyl)-5-phenyl-1H-imidazol-2-yl]-1-methyl-1H-indole;

3-[4,5-bis(4-methoxydiphenyl)-1H-imidazol-2-yl]-1-methyl-1H-indole;

4,4'-[2-(2-phenyl-1H-indol-3-yl)-1H-imidazole-4,5-diyl]bis[N,N-dimethyl]benzenamine;

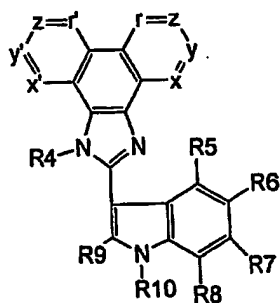
4,4'-[2-(5-chloro-1H-indol-3-yl)-1H-imidazole-4,5-diyl]bis[N,N-dimethyl]benzenamine;

2-(3-indolyl)-4,5-bis[4-(dimethylamino)phenyl]imidazole;

2-(3-indolyl)-4,5-bis[4-(diethylamino)phenyl]imidazole;

2-(2-phenyl-3-indolyl)-4,5-bis[4-dimethylamino)phenyl]imidazole;
2-(2-chloro-3-indolyl)-4,5-bis[4-dimethylamino)phenyl]imidazole;
2-(2-ethylcarboxylate-3-indolyl)-4,5-bis[4-dimethylamino)phenyl]imidazole;
2-(5-chloro-3-indolyl)-4,5-bis[4-(dimethylamino)phenyl]imidazole;
2-(5-cyano-3-indolyl)-4,5-bis[4-(dimethylamino)phenyl]imidazole;
2-(5-nitro-3-indolyl)-4,5-bis[4-(dimethylamino)phenyl]imidazole.

5. A method of inhibiting neoplastic cell growth in a mammal comprising administering to said mammal a therapeutically effective amount of a compound of formula VI:



VI

or a salt thereof, wherein:

R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano;

R10 is H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, acyl;

x is CR11 or N;

y is CR12 or N;

z is CR13 or N;

r is CR14 or N;

x' is CR15 or N;

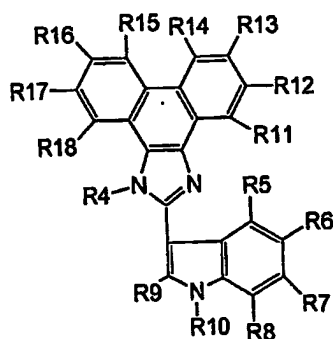
y' is CR16 or N;

z' is CR17 or N;

x' is CR18 or N;

R11, R12, R13, R14, R15, R16, R17 and R18 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, alkenyl, alkenyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano.

6. A method of inhibiting neoplastic cell growth in a mammal comprising administering to said mammal a therapeutically effective amount of a compound of formula VII:



VII

or a salt thereof, wherein:

R4, R5, R6, R7, R8 and R9 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, lower alkenyl, substituted lower alkenyl, lower alkynyl, substituted lower alkynyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano;

R10 is H, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, acyl;

R11, R12, R13, R14, R15, R16, R17 and R18 are independently selected from hydrogen, halogen, hydroxyl, thiol, lower alkyl, substituted lower alkyl, alkenyl, alkenyl, alkylalkenyl, alkyl alkynyl, alkoxy, alkylthio, acyl, aryloxy, amino, amido, carboxyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heterocycle, heteroalkyl, cycloalkyl, substituted cycloalkyl, alkylcycloalkyl, alkylcycloheteroalkyl, nitro, or cyano.

ABSTRACT

The present invention provides therapeutically effective 2,4,5-trisubstituted imidazole compounds, methods of preparing the same, and compositions comprising the compounds alone or in combination with other agents. The present invention further provides for the use of the compounds as anti-cancer agents.

Figure 1:

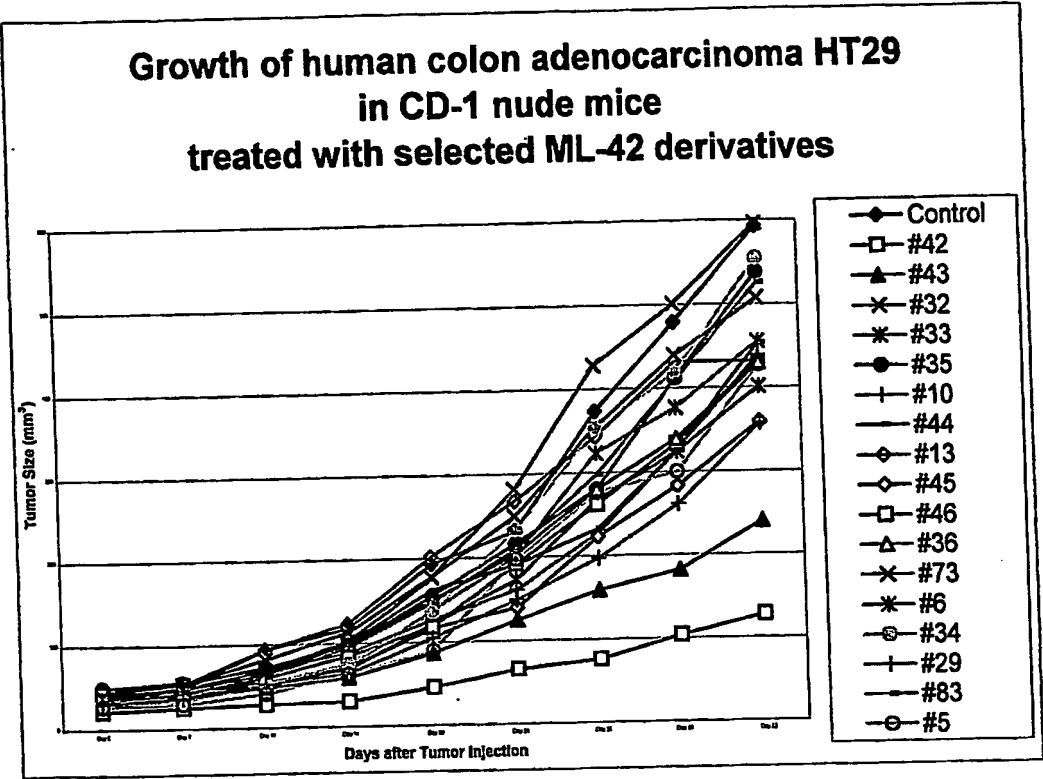


Figure 2

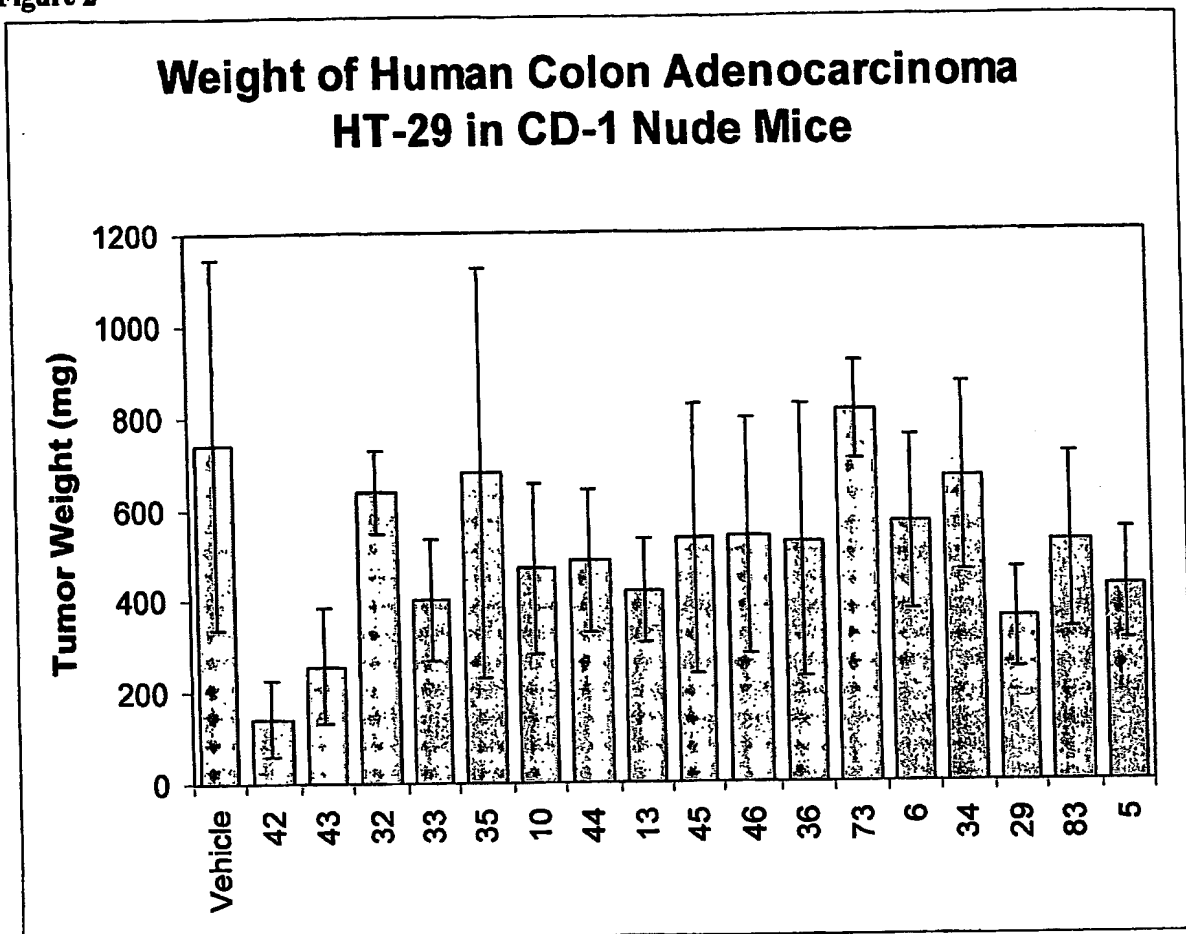
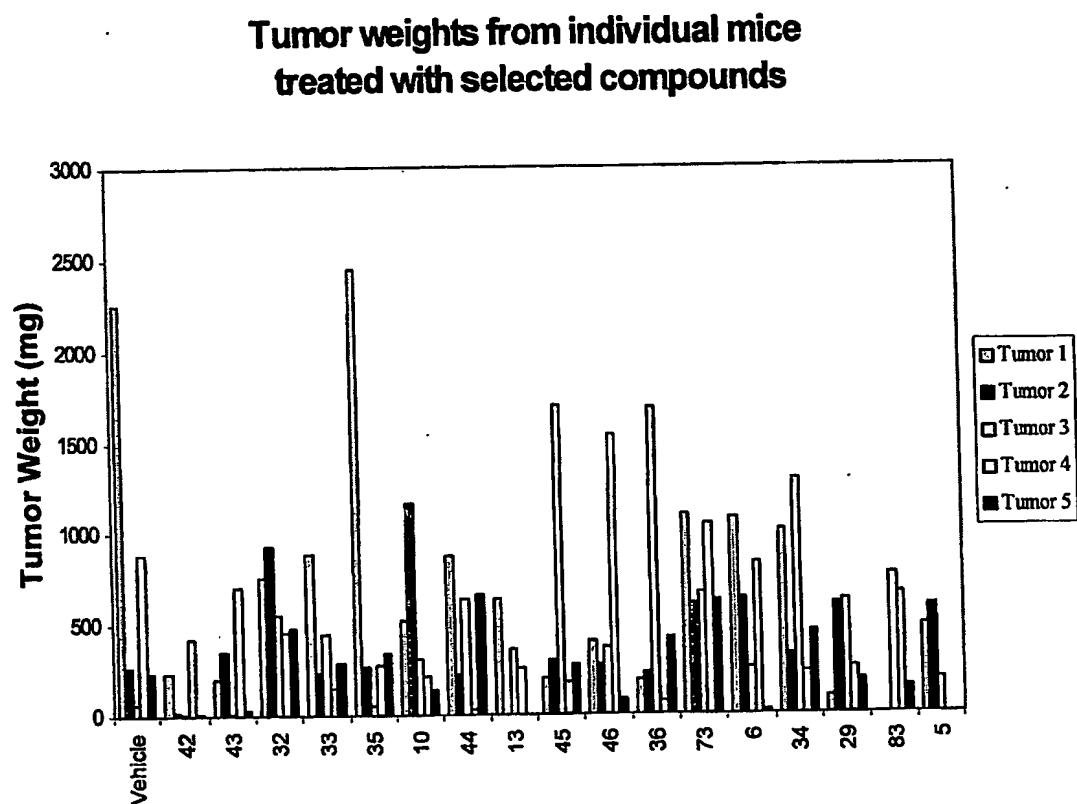


Figure 3



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